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YEAST CELL SURFACE DISPLAY OF PROTEINS AND USES THEREOF

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YEAST CELL SURFACE DISPLAY OF PROTEINS AND USES THEREOF

Cross-Reference to Related Applications

This application is a divisional of U.S. Serial No. 09/724,108, filed November 28, 2000, which is a continuation of U.S. Serial No. 09/009,388, filed January 20, 1998, which is a continuation-in-part of U.S. Serial No. 08/866,398, filed May 30, 1997, now abandoned, which claims benefit of 60/018,741, filed May 31, 1996.

Federal Funding Legend

This invention was produced in part using funds obtained through a grant from the National Institutes of Health grant R01AI35990. Consequently, the federal government has certain rights in this invention.

BACKGROUND OF THE INVENTION

Field of the Invention

The present invention relates generally to the fields of immunology and protein chemistry. More specifically, the present invention relates to the display of peptides and proteins on the yeast cell surface for selection of sequences with desirable binding properties from combinatorial libraries.

Description of the Related Art

Antibody combining site structure can be predicted with reasonable accuracy from polypeptide sequence data, but the ability

to rationally engineer improvements in binding affinity and specificity has proven more elusive, despite some successes (e.g., Roberts et al., '87; Riechmann et al., '92). As a result, mutagenesis and screening of libraries currently represents the most fruitful approach to directed affinity maturation of antibodies. The recent explosion of interest in combinatorial libraries for isolation of molecules with useful binding or catalytic properties has been driven largely by the availability of new techniques for the construction and screening of such libraries. In particular, the construction and screening of antibody immune repertoires *in vitro* promises improved control over the strength and specificity of antibody-antigen interactions.

The most commonly used system for construction of diverse antibody libraries *in vitro* is fusion of antibodies to the coat proteins of filamentous phage (e.g., Huse et al., '89; Clackson et al.,'91; Marks et al.,'92). Fusions are made most commonly to a minor coat protein, called the gene III protein (pIII), which is present in three to five copies at the tip of the phage. A phage constructed in this way can be considered a compact genetic "unit", possessing both the phenotype (binding activity of the displayed antibody) and genotype (the gene coding for that antibody) in one package.

Antibodies possessing desirable binding properties are selected by binding to immobilized antigen in a process called "panning." Phage bearing nonspecific antibodies are removed by washing, and then the bound phage are eluted and amplified by infection of *E. coli.* This approach has been applied to generate antibodies against many antigens, including: hepatitis B surface

antigen (Zebedee et al., '92); polysaccharides (Deng et al., '94), insulin-like growth factor 1 (Garrard & Henner,'93), 2-phenyloxazol-5-one (Riechmann & Weill, '93), and 4-hydroxy-5-iodo-3-nitro-phenacetyl-(NIP)-caproic acid (Hawkins et al., '92).

Although panning of antibody phage display libraries is a powerful technology, it possesses several intrinsic difficulties that limit its wide-spread successful application. First, very high affinity antibodies ($K_D \leq 1$ nM) are difficult to isolate by panning, since the elution conditions required to break a very strong antibody-antigen interaction are generally harsh enough (e.g., low pH, high salt) to denature the phage particle sufficiently to render it non-infective. Secondly, the requirement for physical immobilization of an antigen to a solid surface produces many artifactual difficulties. For example, high antigen surface density introduces avidity effects which mask true affinity. Also, physical tethering reduces the translational and rotational entropy of the antigen, resulting in a smaller ΔS upon antibody binding and a resultant overestimate of binding affinity relative to that for soluble antigen and large effects from variability in mixing and washing procedures lead to difficulties with reproducibility. Thirdly, the presence of only one to a few antibodies per phage particle introduces substantial stochastic variation, and discrimination between antibodies of similar affinity becomes impossible. For example, affinity differences of 6-fold or greater are often required for efficient discrimination (Riechmann & Weill, '93). Finally, populations can be overtaken by more rapidly growing wildtype phage. In particular, since pIII is involved directly in the

phage life cycle, the presence of some antibodies or bound antigens will prevent or retard amplification of the associated phage.

Display of antibodies on the surface of *Escherichia coli* has been developed as an alternative methodology solving several of the problems associated with phage display (Francisco, et al., '93), but introduces new limitations. *E. coli* possesses a lipopolysaccharide layer or capsule that may interfere sterically with macromolecular binding reactions. In fact, a presumed physiological function of the bacterial capsule is restriction of macromolecular diffusion to the cell membrane, in order to shield the cell from the immune system (DiRienzo et al., '78). Since the periplasm of *E. coli* has not evolved as a compartment for the folding and assembly of antibody fragments, expression of antibodies in *E. coli* has typically been very clone dependent, with some clones expressing well and others not at all. Such variability introduces concerns about equivalent representation of all possible sequences in an antibody library expressed on the surface of *E. coli.*

The potential applications of monoclonal antibodies to the diagnosis and treatment of human disease are far-reaching (e.g., Zaccolo & Malavasi, '93; Serafini, '93). Applications to cancer therapy (Hand et al,, '94; Goldenberg, '93; Yarmush et al., '93) and tumor imaging in particular (Fischman et al., '93; Goldenberg & Sharkey, '93; McKearn, '93) have been pursued actively. Antibody therapies for Gram-negative sepsis still hold promise despite discouraging preliminary results (Baumgartner & Glauser, '93). *In vitro* applications to immunohistochemistry (Mietlinen, '93), immunoassay (Kricka, '93; Ishikawa et al., '93), and immunoaffinity

chromatography (Yarmush et al., '92) are already well-developed. For each of these applications, antibodies with high affinity (i.e., $K_D \leq$ 10 nM) and high specificity are desirable. Anecdotal evidence, as well as the *a priori* considerations discussed previously, suggest that phage display is unlikely to consistently produce antibodies of sub-nanomolar affinity.

The structural similarities between B-cells displaying antibodies and yeast cells displaying antibodies provide a closer analogy to *in vivo* affinity maturation than is available with filamentous phage. Moreover, the ease of growth culture and facility of genetic manipulation available with yeast will enable large populations to be mutagenized and screened rapidly. By contrast with conditions in the mammalian body, the physicochemical conditions of binding and selection can be altered for a yeast culture within a broad range of pH, temperature, and ionic strength to provide additional degrees of freedom in antibody engineering experiments.

Combinatorial library screening and selection methods have become a common tool for altering the recognition properties of proteins (Ellman et al., 1997, Phizicky & Fields, 1995). The most widespread technique is phage display, whereby the protein of interest is expressed as a polypeptide fusion to a bacteriophage coat protein and subsequently screened by binding to immobilized or soluble biotinylated ligand. Phage display has been successfully applied to antibodies, DNA binding proteins, protease inhibitors, short peptides, and enzymes (Choo & Klug, 1995, Hoogenboom, 1997, Ladner, 1995, Lowman et al., 1991, Markland et al., 1996, Matthews

& Wells, 1993, Wang et al., 1996). Nevertheless, phage display possesses several shortcomings. For example, some eucaryotic secreted proteins and cell surface proteins require post-translational modifications such as glycosylation or extensive disulfide isomerization which are unavailable in bacterial cells. Furthermore, the nature of phage display precludes quantitative and direct discrimination of ligand binding parameters.

Several bacterial cell surface display methods have been developed (Georgiou et al., 1997). However, use of a procaryotic expression system occasionally introduces unpredictable expression biases (Knappik & Pluckthun, 1995, Ulrich et al., 1995, Walker & Gilbert, 1994) and bacterial capsular polysaccharide layers present a diffusion barrier that restricts such systems to small molecule ligands (Roberts, 1996).

The discovery of novel therapeutics would be facilitated by the development of yeast selection systems. The development of a yeast surface display system for screening combinatorial antibody libraries and a screen based on antibody-antigen dissociation kinetics with the anti-fluorescein scFv-4-4-20 has been described.

The importance of T cell receptors to cell-mediated immunity has been known since the 1980's, but no method for engineering higher affinity T cell receptors has been developed. Although several groups have produced single-chain T cell receptor constructs, these expression systems have allowed biochemical analysis of T cell receptor binding, but have not enabled library methods for altering those binding properties in a directed fashion.

To date, yeast display will fill this gap and as such should be a key technology of tremendous commercial and medical significance.

The prior art is deficient in the lack of effective means of displaying cell surface peptides and proteins for selection of sequences with desirable binding properties. The prior art is also deficient in the lack of effective means of engineering the T cell receptor for improved binding properties. More specifically, no technology has been available to engineer soluble T cell receptors to produce therapeutic intervention of cell-mediated immunity. The present invention fulfills this longstanding need and desire in the art.

SUMMARY OF THE INVENTION

In one embodiment of the present invention, there is provided a genetic method for tethering polypeptides to the yeast cell wall in a form accessible for protein-protein binding. Combining this method with fluorescence-activated cell sorting provides a means of selecting proteins with increased or decreased affinity for another molecule, altered specificity, or conditional binding.

In another embodiment of the present invention, there is provided a method of genetic fusion of a polypeptide of interest to the C-terminus of the yeast Aga2p cell wall protein. Under mating conditions, the outer wall of each yeast cell contains about 10^4 protein molecules called agglutinins. The agglutinins serve as specific contacts to mediate adhesion of yeast cells of opposite mating type during mating. In effect, yeast has evolved a platform for protein-protein binding without steric hindrance from cell wall

components. By attaching an antibody to the agglutinin, one effectively can mimic the cell surface display of antibodies by B cells in the immune system.

In yet another embodiment of the present invention, there is provided a method of fusing a nine residue epitope (HA) tag to the C-terminus of the AGA2 protein. This short peptide is accessible on the cell surface to an antibody in solution without any fixation or digestion of the cells, and can be detected by flow cytometry or fluorescence microscopy. Thus, yeast can be used to display peptides.

In yet another embodiment of the present invention, there is provided a method of fusing an scFv fragment of the 4-4-20 monoclonal antibody to the C-terminus of the AGA2 protein. This fragment is accessible on the cell surface and binds the fluorescein antigen without any fixation or digestion of the cells, and can be detected by flow cytometry or fluorescence microscopy. Thus, yeast can be used to display antibody fragments.

One aspect of the present invention provides a method for selecting proteins with desirable binding properties comprising: transforming yeast cells with a vector expressing a protein to be tested fused at its N-terminus to a yeast cell wall binding protein; labeling the yeast cells with a first label, wherein the first label associates with yeast expressing the protein to be tested and does not associate with yeast which do not express the protein to be tested; selecting for the yeast cells with which said first label is associated; and quantitating said first label, wherein a high occurrence of the first label indicates the protein to be tested has

desirable binding properties and wherein a low occurrence of the first label indicates the protein to be tested does not have desirable binding properties. A preferred embodiment of the present invention further includes the steps of: labeling the yeast cells with a second label, wherein the second label associates with yeast expressing an epitope tag fused to the protein to be tested and encoded by said vector and does not associate with yeast which do not express the epitope tag encoded by said vector; quantitating said second label, wherein an occurrence of the second label indicates a number of expressed copies of the epitope-tagged protein to be tested on the yeast cell surface; and comparing said quantitation of the first label to said quantitation of the second label to determine the occurrence of the first label normalized for the occurrence of the second label, wherein a high occurrence of the first label relative to the occurrence of the second label indicates the protein to be tested has desirable binding properties. Another preferred embodiment of the present invention includes the steps of: labeling the yeast cells with a third label that competes with said first label for binding to the protein to be tested; labeling the yeast cells with said first label; quantitating said first label; labeling the yeast cells with said second label; quantitating said second label; and comparing said quantitation of the first label to said quantitation of the second label to determine the occurrence of the first label normalized for the occurrence of the second label, wherein a low occurrence of the first label relative to the occurrence of the second label indicates the protein to be tested has desirable binding properties.

In one embodiment of the present invention, the first label is a fluorescent label attached to a ligand and the second label is a fluorescent label attached to an antibody. When the labels are fluorescent, the

quantitation step is performed by flow cytometry or confocal fluorescence microscopy.

Another aspect of the present invention provides a vector for performing the method of the present invention, comprising a cell wall binding protein fused to an N-terminus of a protein of interest. Preferred embodiments of this aspect of the present invention include means for expressing a polypeptide epitope tag fused to said protein of interest in said yeast cells. A more preferred embodiment provides that the cell wall binding protein is the binding subunit of a yeast agglutinin protein, even more preferably yeast agglutinin binding subunit is Aga2p.

Another preferred embodiment of the present aspect of the invention provides that the epitope tag amino acid sequence is selected from the group of YPYDVPDYA (HA), EQKLISEEDL (c-*myc*), DTYRYI, TDFYLK, EEEEYMPME, KPPTPPPEPET, HHHHHH, RYIRS, or DYKDDDDK, and that the N-terminus of said protein of interest is fused to a C-terminus of said cell wall binding protein.

Yeast surface display and sorting by flow cytometry have been used to isolate mutants of a scFv that is specific for the Vb8 region of the T cell receptor. Selection was based on equilibrium binding by two fluorescently-labeled probes, a soluble Vb8 domain and an antibody to the c-*myc* epitope tag present at the carboxy-terminus of the scFv. The mutants that were selected in this screen included a scFv with three-fold increased affinity for the Vb8 and scFv clones that were bound with reduced affinities by the anti-c-*myc* antibody. The latter finding indicates that the yeast display system may be used to map conformational epitopes, which can not

be revealed by standard peptide screens. Equilibrium antigen binding constants were estimated within the surface display format, allowing screening of isolated mutants without necessitating subcloning and soluble expression. Only a relatively small library of yeast cells (3×10^5) displaying randomly mutagenized scFv was screened to identify these mutants, indicating that this system will provide a powerful tool for engineering the binding properties of eucaryotic secreted and cell surface proteins.

Another preferred embodiment of the present aspect of the invention provides a method for displaying proteins that are not displayed as their normal ("wild type") sequence. In the example shown, the T cell receptor for antigen was not expressed as its "wild type" sequence. However, after random mutagenesis and selection by flow cytometry with appropriate conformationally-specific antibodies, the mutant receptors were expressed on the yeast cell surface. This strategy will allow the discovery of novel T cell receptors and it provides a method for the display of virtually any polypeptide. Thus, the present invention also provides a method for selecting proteins for displayability on a yeast cell surface, comprising the step of: transforming yeast cells with a vector expressing a protein to be tested fused to a yeast cell wall protein, wherein mutagenesis is used to a generate a variegated population of mutants of the protein to be tested; labeling said yeast cells with a first label, wherein said first label associates with yeast expressing said protein to be tested and does not associate with yeast which do not express said protein to be tested; isolating said yeast cells with which said first label is associated, by quantitating said first label,

wherein a high occurrence of said first label indicates said protein to be tested has desirable display properties and wherein a low occurrence of said first label indicates said protein to be tested does not have desirable display properties. Preferably, the protein tested is an antibody, Fab, Fv, or scFv antibody fragment or the ligand binding domain of a cell surface receptor. A representative example of a cell surface receptor is a T cell receptor.

Other and further aspects, features, and advantages of the present invention will be apparent from the following description of the presently preferred embodiments of the invention given for the purpose of disclosure.

BRIEF DESCRIPTION OF THE DRAWINGS

So that the matter in which the above-recited features, advantages and objects of the invention are attained and can be understood in detail, more particular descriptions of the invention may be had by reference to certain embodiments thereof which are illustrated in the appended drawings. These drawings form a part of the specification. It is to be noted, however, that the appended drawings illustrate preferred embodiments of the invention and therefore are not to be considered limiting in their scope.

Figure 1 is a schematic, showing *in vitro* affinity maturation by yeast display.

Figure 2 shows the schematic illustration of surface display on yeast. A nine amino acid peptide epitope from the hemagglutinin antigen (HA) was fused to the C-terminus of the Aga2p subunit of a-agglutinin.

followed by the 4-4-20 anti-fluorescein scFv sequence. An additional ten residue epitope tag (c-*myc*) was fused at the C-terminus of the scFv, allowing quantitation of fusion display independent of antigen binding by either the HA or c-myc tags. The HA or c-*myc* tag can be used to normalize for variation in the number of displayed fusion proteins in double-label flow cytometry.

Figure 3 shows a vector for yeast surface display. Figure 3A shows the construction of the vector pCT202. Figure 3B shows the specific restriction sites and illustrates the transcriptional regulation by galactose, the N-terminal HA and C-terminal c-*myc* epitope tags and the Factor Xa protease cleavage site.

Figure 4 demonstrates that the displayed fusions can be detected by fluorescence techniques, showing a flow cytometric histogram of yeast labeled with α-c-*myc*/α-mouse-PE.

Figure 5 demonstrates that antigen binding by 4-4-20 scFv can be detected by fluorescence, showing a flow cytometric histogram of yeast labeled with FITC-dextran (2 x 10^6 Da).

Figure 6 shows that 4-4-20 activity and c-*myc* can be detected simultaneously, and demonstrate a 1:1 correlation of fluorescence signals; therefore, variation in intensity signal 1 (FITC) can be normalized for cell-to-cell variation in expression of the protein of interest by the intensity of signal 2 (PE).

Figure 7 shows the sequence of the AGA2-HA-4-4-20-c-myc gene cassette.

Figure 8 shows confocal microscopic images of yeast displaying scFv. Yeast containing plasmid directing surface

expression of the HA peptide (**Figure 8A**) or the scFv fusion (**Figure 8B**) were labeled with mAb 9E10, followed by a secondary anti-mouse IgG-R-phycoerythrin (PE) conjugate and FITC-dextran. DIC (upper panels), red PE fluorescence (middle panels), and green FITC fluorescence (lower panels) images were collected.

Figure 9 shows flow cytometric analyses of yeast displaying scFv. Yeast strains displaying either (**Figure 9A**) an irrelevant peptide or (**Figure 9B**) the 4-4-20 scFv were labeled with mAb 9E10 and FITC-dextran. Cells displaying scFv were also treated with 5 mM DTT prior to labeling (**Figure 9C**). (i) Univariate histograms of PE fluorescence associated with labeling by 9E10; (ii) univariate histograms of FITC fluorescence; (iii) bivariate histograms showing correlation between PE and FITC fluorescence.

Figure 10 demonstrates the enrichment of yeast displaying improved scFv variants by kinetic selection and flow cytometric cell sorting. Yeast expressing a mutagenized 4-4-20 scFv library (**Figure 10A**) and a yeast pool resulting from three rounds of kinetic selection and amplification (**Figure 10B**) were subjected to competitive dissociation of fluorescent antigen with 5-aminofluorescein, leaving cells displaying the tightest binding mutants with the highest ratio of FITC intensity/PE intensity.

Figure 11 shows dissociation kinetics of the interaction between fluorescein and surface displayed scFv. Yeast displaying 4-4-20 scFv (circles), mutant 4M1.1 (squares) isolated from the library, and mutant 4M1.2 (triangles) were labeled with mAb 9E10 and FITC-dextran. 5-aminofluorescein was added as a competitor. Mean

intensity of FITC fluorescence of the 9E10 positive population of cells was followed as a function of time. The slope of the line is equal to the kinetic dissociation rate, and the extrapolated value at time t = 0 sec is equal to the valency of the interaction. MFI_i = relative mean fluorescence intensity of yeast at time t = i.

Figure 12 shows the expression levels and antigen binding properties of yeast surface displayed scFv-KJ16 (shaded) and control Aga2p/HA (unshaded). Yeast strain EBY100 was transformed with scFv-KJ16 cloned into the yeast display vector pCT202 or the pCT202 vector alone. After induction in galactose medium at 20°C overnight, cells were stained with fluorescent antibodies and analyzed by flow cytometry. (**Figure 12A**) scFv-KJ16/yeast or Aga2p/HA/yeast stained with mouse anti-HA Mab (12CA5) followed by FITC-labeled goat anti-mouse IgG, (**Figure 12B**) scFv-KJ16/yeast or Aga2p/HA/yeast stained with mouse anti-c-*myc* Mab (9E10) followed by FITC-labeled goat anti-mouse IgG, (**Figure 12C**) scFv-KJ16/yeast or Aga2p/HA/yeast stained with biotinylated-scTCR at ~10 nM followed by a streptavidin-phycoerythrin conjugate, and (**Figure 12D**) scFv-KJ16/yeast stained with biotinylated-scTCR followed by a streptavidin-phycoerythrin conjugate in the presence (shaded) or absence (unshaded) of intact IgG KJ16 at 100 mg/ml.

Figure 13 shows the equilibrium antigen binding isotherm of cell wall displayed scFv-KJ16, determined by flow cytometry. Yeast strain EBY100 displaying surface scFv-KJ16 was incubated with varying concentrations of biotinylated-scTCR, labeled with a streptavidin-phycoerythrin conjugate, and detected by flow cytometry. Data was plotted as a Scatchard diagram or as a titration

(inset) and an effective K_D ~500 nM was determined. MFU refers to mean fluorescence units.

Figure 14 shows the two dimensional fluorescence histograms and sorting window used to select scFv-KJ16 mutants. scFv-KJ16 cloned into the display vector pCT202 was transformed into the *E. coli* mutator strain XL1-Red (Stratagene) and propagated for six overnight growth cycles. Plasmids of the mutant library were purified and used in LiAc transformation (Gietz et al., 1995) of EBY100 yeast. After induction at 30°C, yeast were sorted using a fluorescence-activated cell sorter. (**Figure 14A**) Representative histogram from the first round of cell sorting, with the sorting window indicated, and (**Figure 14B**) representative histogram from the fourth (final) round of sorting, illustrating an enrichment of the population in the sorting window.

Figure 15 shows the mean levels of binding to anti-HA Mab, anti-c-*myc* Mab, or biotinylated-scTCR for ten randomly selected clones from the final sort shown in Figure 14B. Ten mutants and wt scFv-KJ16/yeast were induced in galactose medium at 30°C overnight. Cells were analyzed by flow cytometry after staining with mouse anti-HA Mab followed by FITC-labeled goat anti-mouse IgG (open bars), mouse anti-c-*myc* followed by FITC-labeled goat anti-mouse IgG (gray bars), or biotinylated-scTCR at ~40 nM followed by a streptavidin-phycoerythrin conjugate (black bars).

Figure 16 shows the fluorescent labeling distributions for anti-c-*myc* or scTCR binding of three selected mutants shown in Figure 4. Three classes of scFv-KJ16/yeast mutants were double-

stained with anti-c-*myc* and biotinylated-scTCR followed by FITC-labeled goat anti-mouse IgG and a streptavidin-phycoerythrin conjugate, then analyzed by flow cytometry as described in Figure 4. The fluorescent distributions for each scFv-KJ16/yeast mutant (shaded) and wt scFv-KJ16/yeast (unshaded) are shown. **Figures 16A and 16B**, mut4; **Figures 16C and 16D**, mut7; **Figures 16E and 16F**, mut10.

Figure 17 shows the equilibrium antigen binding isotherms for three mutants shown in Figure 16. Aga2p/HA/yeast, wt scFv-KJ16/yeast, and three mutant scFv-KJ16/yeast characterized in Figure 16 were stained with various dilutions of biotinylated-scTCR followed by a streptavidin-phycoerythrin conjugate. After analysis by flow cytometry, binding isotherms were graphed with MFU as a function of scTCR dilution.

Figure 18 shows the sequence analysis of wild-type scFv-KJ16, mut4, and mut7. Plasmids from wt scFv-KJ16/yeast and two mutants (mut4 and mut7) were recovered by plasmid rescue and transformed into *E. coli* DH5a competent cells to produce plasmids for sequencing, as described below. Sequence analysis was performed using primers that flank the scFv of the display vector. Mutations are indicated in bold.

Figure 19 shows the flow cytometry profiles of antibody binding to yeast that have been transformed with a plasmid that contains a T cell receptor single-chain (VαVβ) gene. The normal or wild type (wt) sequence is compared with several mutants (mTCR7, mTCR15, mTCR16) that were selected after random mutagenesis of the scTCR plasmid. Selection involved binding of the antibody 1B2,

which recognizes a conformational epitope on the T cell receptor, followed by several round of fluorescent-activated cell sorting. In the first panel, the yeast cells were stained with an antibody (12CA5) to the HA tag. In the second panel, the yeast cells were stained with an antibody (1B2) to the T cell receptor. Although the HA epitope is expressed on the surface in each case, only those cells that express a mutagenized plasmid are capable of expressing the native T cell receptor (1B2 positive).

Figure 20 shows the flow cytometry profiles of antibody binding to yeast that have been transformed with double mutants from the selection shown in Figure 19. Cells were stained for flow cytometry as described in Figure 19. Double mutants expressed an increase in level of the T cell receptor (i.e. 1B2-reactive material). The results show that by combining single mutations it is possible to enhance the level of cell surface expression of the T cell receptor.

Figure 21 shows the sequence of mutations that lead to the enhanced expression of the cell surface T cell receptor. These included residues 17 of the Vβ, 43 of the Vα, and 104 of the Vα.

DETAILED DESCRIPTION OF THE INVENTION

As used herein, the term "affinity maturation" shall refer to a process of successive mutation and selection by which antibodies of higher affinity are selected.

As used herein, the term "agglutinin" shall refer to a yeast surface adhesion protein which binds two cells together during mating.

As used herein, the term "antibody" shall refer to a protein produced by mammalian immune systems which binds tightly and specifically to particular molecules.

As used herein, the term "ligand" shall refer to a molecule that is bound specifically by a particular protein.

As used herein, the term "antigen" shall refer to a ligand that is bound specifically by an antibody.

As used herein, the term "Complementarity Determining Region" or "CDR" shall refer to the portion of an antibody which directly contacts the bound antigen.

As used herein, the term "Fluorescence Activated Cell Sorting" or "flow cytometry" shall refer to a method for sorting cell populations on the basis of differential fluorescent labeling.

As used herein, the term "hapten" shall refer to a small antigen which cannot stimulate an immune response without being conjugated to a carrier.

As used herein, the term "single chain antibody" or "SCA" shall refer to a fusion of portions of the heavy and light chains of an antibody which retains a single active binding site. The term scFv is used interchangeably to refer to a single chain antibody.

As used herein, the term "epitope tag" shall refer to a contiguous sequence of amino acids specifically bound by an antibody when fused to another protein.

As used herein, the term "HA" refers to the epitope tag sequence YPYDVPDYA.

As used herein, the term "c-*myc*" refers to the epitope tag sequence EQKLISEEDL.

As used herein, the term "scFv 4-4-20" refers to an scFv which binds specifically to fluorescein and fluorescein conjugated to other molecules such as biotin or dextran.

As used herein, the term "AGA2p" refers to the protein product of the yeast AGA2 mating type a agglutinin gene.

The term "displayability" will be used to describe a combination of biophysical characteristics allowing a protein to escape the secretory "quality control" apparatus that retains and degrades misfolded proteins (Hammond & Helenius, 1995.) Proteins displayed on the yeast cell surface must first pass successfully through the quality control step. Protein folding kinetics and thermodynamic stability together are believed to determine the efficiency of escape from the quality control apparatus.

In accordance with the present invention there may be employed conventional molecular biology, microbiology, and recombinant DNA techniques within the skill of the art. Such techniques are explained fully in the literature. See, e.g., Maniatis, Fritsch & Sambrook, "Molecular Cloning: A Laboratory Manual (1982); "DNA Cloning: A Practical Approach," Volumes I and II (D.N. Glover ed. 1985); "Oligonucleotide Synthesis" (M.J. Gait ed. 1984); "Nucleic Acid Hybridization" (B.D. Hames & S.J. Higgins eds. (1985)); "Transcription and Translation" (B.D. Hames & S.J. Higgins eds. (1984)); "Animal Cell Culture" (R.I. Freshney, ed. (1986));

"Immobilized Cells And Enzymes" (IRL Press, (1986)); B. Perbal, "A Practical Guide To Molecular Cloning" (1984).

Therefore, if appearing herein, the following terms shall have the definitions set out below.

A "vector" is a replicon, such as plasmid, phage or cosmid, to which another DNA segment may be attached so as to bring about the replication of the attached segment

A "DNA molecule" refers to the polymeric form of deoxyribonucleotides (adenine, guanine, thymine, or cytosine) in either single stranded form, or a double-stranded helix. This term refers only to the primary and secondary structure of the molecule, and does not limit it to any particular tertiary forms. Thus, this term includes double-stranded DNA found, inter alia, in linear DNA molecules (e.g., restriction fragments), viruses, plasmids, and chromosomes. In discussing the structure herein according to the normal convention of giving only the sequence in the 5' to 3' direction along the nontranscribed strand of DNA (i.e., the strand having a sequence homologous to the mRNA).

A DNA "coding sequence" is a double-stranded DNA sequence which is transcribed and translated into a polypeptide *in vivo* when placed under the control of appropriate regulatory sequences. The boundaries of the coding sequence are determined by a start codon at the 5' (amino) terminus and a translation stop codon at the 3' (carboxyl) terminus. A coding sequence can include, but is not limited to, prokaryotic sequences, cDNA from eukaryotic mRNA, genomic DNA sequences from eukaryotic (e.g., mammalian) DNA, and even synthetic DNA sequences. A polyadenylation signal

and transcription termination sequence will usually be located 3' to the coding sequence.

Transcriptional and translational control sequences are DNA regulatory sequences, such as promoters, enhancers, polyadenylation signals, terminators, and the like, that provide for the expression of a coding sequence in a host cell.

A "promoter sequence" is a DNA regulatory region capable of binding RNA polymerase in a cell and initiating transcription of a downstream (3' direction) coding sequence. For purposes of defining the present invention, the promoter sequence is bounded at its 3' terminus by the transcription initiation site and extends upstream (5' direction) to include the minimum number of bases or elements necessary to initiate transcription at levels detectable above background. Within the promoter sequence will be found a transcription initiation site (conveniently defined by mapping with nuclease S1), as well as protein binding domains (consensus sequences) responsible for the binding of RNA polymerase. Eukaryotic promoters will often, but not always, contain "TATA" boxes and "CAT" boxes. Prokaryotic promoters contain Shine-Dalgarno sequences in addition to the -10 and -35 consensus sequences.

An "expression control sequence" is a DNA sequence that controls and regulates the transcription and translation of another DNA sequence. A coding sequence is "under the control" of transcriptional and translational control sequences in a cell when RNA polymerase transcribes the coding sequence into mRNA, which is then translated into the protein encoded by the coding sequence.

A "selection gene" refers to a gene that enables the discrimination of cells displaying a required phenotype upon implementation of certain conditions. For example, the growth of bacteria in medium containing antibiotics to select for the bacterial cells containing antibiotic resistance genes.

The term "primer" as used herein refers to an oligonucleotide, whether occurring naturally as in a purified restriction digest or produced synthetically, which is capable of acting as a point of initiation of synthesis when placed under conditions in which synthesis of a primer extension product, which is complementary to a nucleic acid strand, is induced, i.e., in the presence of nucleotides and an inducing agent such as a DNA polymerase and at a suitable temperature and pH. The primer may be either single-stranded or double-stranded and must be sufficiently long to prime the synthesis of the desired extension product in the presence of the inducing agent. The exact length of the primer will depend upon many factors, including temperature, the source of primer and the method used. For example, for diagnostic applications, depending on the complexity of the target sequence, the oligonucleotide primer typically contains 15-25 or more nucleotides, although it may contain fewer nucleotides.

The primers herein are selected to be "substantially" complementary to different strands of a particular target DNA sequence. This means that the primers must be sufficiently complementary to hybridize with their respective strands. Therefore, the primer sequence need not reflect the exact sequence of the template. For example, a non-complementary nucleotide

fragment may be attached to the 5' end of the primer, with the remainder of the primer sequence being complementary to the strand. Alternatively, non-complementary bases or longer sequences can be interspersed into the primer, provided that the primer sequence has sufficient complementarity with the sequence or hybridize therewith and thereby form the template for the synthesis of the extension product.

As used herein, the terms "restriction endonucleases" and "restriction enzymes" refer to bacterial enzymes, each of which cut double-stranded DNA at or near a specific nucleotide sequence.

A cell has been "transformed" by exogenous or heterologous DNA when such DNA has been introduced inside the cell. The transforming DNA may or may not be integrated (covalently linked) into the genome of the cell. In prokaryotes, yeast, and mammalian cells for example, the transforming DNA may be maintained on an episomal element such as a plasmid. With respect to eukaryotic cells, a stably transformed cell is one in which the transforming DNA has become integrated into a chromosome so that it is inherited by daughter cells through chromosome replication. This stability is demonstrated by the ability of the eukaryotic cell to establish cell lines or clones comprised of a population of daughter cells containing the transforming DNA. A "clone" is a population of cells derived from a single cell or common ancestor by mitosis. A "cell line" is a clone of a primary cell that is capable of stable growth in vitro for many generations.

A "heterologous" region of the DNA construct is an identifiable segment of DNA within a larger DNA molecule that is not

found in association with the larger molecule in nature. Thus, when the heterologous region encodes a mammalian gene, the gene will usually be flanked by DNA that does not flank the mammalian genomic DNA in the genome of the source organism. In another example, coding sequence is a construct where the coding sequence itself is not found in nature (e.g., a cDNA where the genomic coding sequence contains introns, or synthetic sequences having codons different than the native gene). Allelic variations or naturally-occurring mutational events do not give rise to a heterologous region of DNA as defined herein.

As used herein, the term "reporter gene" refers to a coding sequence attached to heterologous promoter or enhancer elements and whose product is easily and quantifiably assayed when the construct is introduced into tissues or cells.

Transcriptional and translational control sequences are DNA regulatory sequences, such as promoters, enhancers, polyadenylation signals, terminators, and the like, that provide for the expression of a coding sequence in a host cell.

A number of fluorescent materials are known and can be utilized as labels. These include, for example, fluorescein, rhodamine, auramine, Texas Red, AMCA blue, R-phycoerythrin, B-phycoerythrin, and Lucifer Yellow.

A number of polypeptide sequences that can be fused to proteins and bound specifically by antibodies are known and can be utilized as epitope tags. These include, for example, HA, c-*myc*, DTYRYI, TDFYLK, EEEEYMPME, KPPTPPPEPET, HHHHHH, RYIRS, and DYKDDDDK.

Antibodies are protein molecules produced by the human immune system to recognize, bind to, and mediate the clearance of foreign substances from the body. Technologies have been developed to take advantage of antibodies for highly-specific cancer diagnosis and therapy. For example, by tethering radioisotopes or toxins to an antibody which binds to tumor cells, it is possible to deliver a focused dosage of such cell-killing agents to the diseased tissue while leaving surrounding tissue comparatively unharmed. Antibodies are also critical tools in biotechnology, and are used extensively for analytical purposes, e.g., to quantify trace quantities of substances and separations, and to purify desired biological products from complex mixtures.

In these applications, both the strength of the antibody bond with its target (affinity) and the selectivity with which an antibody binds to only its particular target (specificity) are crucial. For this reason, protein engineers seek to alter and improve the binding characteristics of particular antibodies. Rational approaches to antibody structural design have met with limited success, and available methods for random screening possess significant limitations.

The mammalian immune system's approach to the problem of fine tuning antibody affinity is by a process called "affinity maturation," wherein cycles of mutation and evolutionary selection produce antibodies which bind their targets more tightly. The present invention discloses a powerful new system for engineering antibody affinity and specificity, by constructing a microbial analog of the mammalian immune system's B cell

repertoire. Antibodies were displayed on the surface of yeast cells by genetic fusion with cell wall proteins. After mutation, variants were selected on the basis of improved binding characteristics with fluorescently labeled targets.

The yeast antibody display method was tested by studying model antibodies whose physical and chemical properties are already well characterized. These methods are then straightforwardly applied to antibodies of practical interest. The genetic malleability of yeast, the ease of growth of this microbe, and the ability to modify antibody binding conditions in the test tube combine to produce unprecedented control over the engineering of antibody affinity and specificity.

The advantage of the library method of the present invention is that it is particularly suited for proteins such as antibodies. The most widely used method currently consists of "panning" for antibodies displayed on the surface of bacteriophage. Yeast display has several advantages over phage display. First, the antibody-antigen bond need not be broken to recover tightly-bound variants. The harsh conditions required for disrupting this bond in prior art methods can reduce infectivity of phage. Secondly, increased library diversity due to decreased clonal deletion is an advantage. It is well known that many antibody structures cannot be correctly processed by the bacterial secretory apparatus. Yeast cells are eucaryotic and possess secretory pathways very similar to mammalian cells. Thirdly, the present invention provides a more accurate and precise determination of antibody-antigen affinity. The presence of 10^4 molecules per cell eliminates the stochastic variation

that results with only a few molecules per phage. Finally, quantitation of fluorescence by flow cytometry provides a continuous measure of surface-bound antigen without *a priori* knowledge of affinity in by comparison to the binary bound/released dichotomy with panning of phage. Also, bacteria possess a lipopolysaccharide layer which acts as a macromolecular permeability barrier preventing antibody or protein access to displayed molecules.

The present invention discloses a surface display system for the *in vitro* expression and selection of peptide and protein libraries on yeast. A nine residue peptide epitope (HA) has been fused to the binding subunit of a yeast cell wall protein (AGA2), followed by the 4-4-20 anti-fluorescein single-chain Fv. Selection was performed by flow cytometry on mixtures of cells with and without the displayed fusion. 600-fold enrichments were achieved in one pass of sorting. The system of the present invention illustrates a process for the *in vitro* affinity maturation of antibodies as well as a process for the directed evolution of other proteins and peptides, with the advantages of (i) a double-label flow cytometry selection scheme allowing finer affinity discrimination than panning; (ii) as many as 10^4 copies of the displayed sequence per cell, eliminating stochastic variations in the selection; and (iii) library expression in yeast, with an altered or potentially improved expression bias which could yield clones that would be deleted from a library expressed in *E. coli.*

One object of the present invention is the engineering of antibodies for improved affinity and specificity. Toward this end, antibody-hapten binding was studied via mutagenesis and screening

of antibodies expressed on the external cell wall of the yeast *Saccharomyces cerevisiae.* As an experimentally facile and genetically pliable eucaryote, yeast presents significant advantages over filamentous phage display as a platform for antibody expression and engineering. In essence, a microbial analog of the mammalian immune system B-cell repertoire was constructed *in vitro,* allowing antibody affinity maturation to be performed under strictly controlled conditions of mutagenesis and selection. As a result, antibodies of significantly improved affinity and specificity were attainable.

One aspect of the present invention provides a method for selecting proteins with desirable binding properties comprising: transforming yeast cells with a vector expressing a protein to be tested fused at its N-terminus to a yeast cell wall binding protein; labeling the yeast cells with a first label, wherein the first label associates with yeast expressing the protein to be tested and does not associate with yeast which do not express the protein to be tested; selecting for the yeast cells with which said first label is associated; quantitating said first label, wherein a high occurrence of the first label indicates the protein to be tested has desirable binding properties and wherein a low occurrence of the first label indicates the protein to be tested does not have desirable binding properties. A preferred embodiment of the present invention further includes the steps of: labeling the yeast cells with a second label, wherein the second label associates with yeast expressing an epitope tag fused to the protein to be tested and encoded by said vector and does not associate with yeast which do not express the epitope tag encoded by said vector;

quantitating said second label, wherein an occurrence of the second label indicates a number of expressed copies of the epitope tagged protein to be tested on the yeast cell surface: and comparing said quantitation of the first label to said quantitation of the second label to determine the occurrence of the first label normalized for the occurrence of the second label. wherein a high occurrence of the first label relative to the occurrence of the second label indicates the protein to be tested has desirable binding properties. Another preferred embodiment of the present invention includes the steps of: labeling the yeast cells with a third label that competes with said first label for binding to the protein to be tested: labeling the yeast cells with said first label; quantitating said first label; labeling the yeast cells with said second label; quantitating said second label: and comparing said quantitation of the first label to said quantitation of the second label to determine the occurrence of the first label normalized for the occurrence of the second label. wherein a low occurrence of the first label relative to the occurrence of the second label indicates the protein to be tested has desirable binding properties.

Another aspect of the present invention provides a vector for performing the method of the present invention, comprising a cell wall binding protein fused to an N-terminus of a protein of interest. Preferred embodiments of this aspect of the present invention include means for expressing a polypeptide epitope tag in said yeast cells. A more preferred embodiment provides that the cell wall binding protein is a yeast agglutinin protein binding subunit, even more preferably yeast agglutinin protein is Aga2p.

The following examples are given for the purpose of illustrating various embodiments of the invention and are not meant to limit the present invention in any fashion.

EXAMPLE 1

Media/Buffers

The following media/buffers were used herein:

E. coli

LB media (1X): Bacto tryptone (Difco, Detroit, MI): 10.0 g; Bacto yeast extract (Difco): 5.0 g; NaCl:10.0 g. Make up to 1 L, autoclave. For plates, add 15g/L Agar and autoclave.

Ampicillin

Stock: 25 mg/ml of sodium salt of ampicillin in water. Filter sterilize and store in aliquots of 4 mls at -20°C. Working [ ] = 35-50 µg/ml; 4 ml of aliquot in 1 L —> 100 µg/ml; 2 ml of aliquot in 1 L —> 50 µg/ml. Add to autoclaved LB only after it has cooled to ~ 55°C.

SOC media 100 mL

2% Bacto tryptone: 2.0 g; 0.5% Yeast Extract (Difco):0.5 g; 10 mM NaCl: 0.2 ml 5 M; 10 mM $MgCl_2$: 1.0 ml 1 M; 10 mM $MgSO_4$: 1.0 ml 1 M; 20 mM Dextrose: 0.36 g. Autoclave or filter sterilize.

YEAST

Synthetic Minimal + Casamino acids (SD-CAA) 500 mL

Dextrose(Glucose) 10.00 g

Yeast Nitrogen Base w/o Amino Acids (Difco)	3.35 g
$Na_2HPO_4 \cdot 7H_2O$	5.1 g
$NaH_2PO_4 \cdot H_2O$	4.28 g
Casamino Acids (Trp-, Ura-) (Difco)	2.5 g

Add dH_2O to final volume. Filter sterilize and refrigerate.

For plates, dissolve sodium phosphates and sorbitol to 1 M final concentration in 400 ml dH_2O. Add 7.5 g agar and autoclave. Dissolve dextrose, N_2 base, and amino acids in 100 ml dH_2O and filter sterilize. Add the filtered reagents after the autoclaved salts have cooled enough to touch.

SG-CAA (Induction Medium)	500 mL
Galactose	10.00 g
Yeast Nitrogen Base w/o Amino Acids (Difco)	3.35 g
$Na_2HPO_4 \cdot 7H_2O$	5.1 g
$NaH_2PO_4 \cdot H_2O$	4.28 g
Casamino Acids (Trp-, Ura-) (Difco)	2.5 g

Add dH_2O to final volume. Filter sterilize and refrigerate.

Rich (YPD) 1000 mL

Yeast extract:10 g; Peptone (Difco):20 g; Dextrose: 20 g. Add dH_2O to 1 L. Autoclave.

TAE (Tris-Acetate)

Working soln: 0.04 M Tris-acetate and 0.001 M EDTA

Stock (50X):	in 1 L
Tris base	242 g

glacial acetic acid	57.1 ml
0.5 M EDTA (pH 8.0)	100 ml

TBE (Tris-Borate): Working solution: 0.09 M Tris-borate and 0.001M EDTA

Stock (5X):	in 1 L
Tris base	54 g
boric acid	27.5 g
0.5 M EDTA (pH 8.0)	20.0 ml

Stop buffer 10X (Restriction):

50% v/v glycerol; 0.1 M EDTA (pH 7.5); 1% w/v SDS; 0.1% w/v bromophenol blue. Combine all components except for the dye and pH to 7.5 before dye addition.

Staining -- Ethidium bromide

0.5 µg/ml in water; Stock: 10 mg/ml. Dilutions of Stock: 1/10 in TBE. Add 100 µl dilution in 100 ml buffer.

TBS Working solution: 10 mM Tris-HCl, 140 mM NaCl and 5 mM EDTA. Filter sterilize.

EXAMPLE 2

Protocol: Replica Plating

1. Choose a material suitable for colony lifts, making sure it is washed, dried and sterile.

2. Mark the bottom of each fresh replica plate with an arrow to line up the plate. Also mark the starting plate.

3. Take the top off the starting plate, turn it upside down and line up the arrow on the bottom with the mark on the colony lift material. Lay the plate down onto the surface of the material and gently put pressure on the entire plate. Make sure the plate doesn't move around after it has touched the material. Remove the plate and replace the lid. Portions of the colonies that transferred to the material can be seen.

4. Repeat this procedure with one of the fresh replica plates. Make sure the arrow lines up with the mark also to make an exact replica. Hold up to the light to see the tiny colonies that transferred.

5. Repeat the entire procedure for each replica plate to be made.

6. Incubate the replica plates at the appropriate conditions for selective growth. Colonies will usually grow up within a day or so.

EXAMPLE 3

Protocol: Electrotransformation of Yeast

The cell preparation procedure was as follows: Step1. Inoculate 50 ml of YPD with an overnight culture to an OD of 0.1. Step 2. Grow cells at 30°C with vigorous shaking to an OD_{600} of 1.3 to

1.5 (approximately 6 hours). Step 3. Harvest in cold rotor at 3500 rpm for 5 minutes at 4°C. Discard supernatant. Step 4. Thoroughly wash the cells by resuspending in 50 ml cold sterile water. Centrifuge as above and discard supernatant. Step 5. Repeat step 4 with 25 ml cold water. Step 6. Resuspend in 2 ml of ice-cold sterile 1 M sorbitol. Centrifuge as above and discard supernatant. Step 7. Resuspend in 50 ml ice-cold 1 M sorbitol. Final volume of cells is about 150 ml (enough for 3 transformations).

Electrotransformation:

1. Place 0.2 cm cuvettes and white slide chamber on ice.

2. In an eppendorf tube, add 50 ml of yeast suspension and gently mix in < 5 ml (0.1 mg) of plasmid DNA in TE. Make sure to add DNA to yeast already in eppendorf. Place on ice for 5 minutes (This time frame is pretty critical).

3. Set GENE PULSER at 1.5 kV and 25 mF. Set the Pulse Controller to 200 W. The time constant for this pulse should be 4.5 to 5.0 msec.

4. Transfer 40 ml of cell/DNA mixture to pre-chilled electroporation cuvette. Tap contents to bottom, making sure the sample is in contact with both aluminum sides of the cuvette. Place the cuvette in chilled safety chamber slide. Push slide into the chamber until the cuvette makes contact with the electrodes in the base of the chamber.

5. Apply one pulse at the settings above.

6. Remove the slide with the cuvette, and immediately add 1 ml of cold 1M sorbitol to the cuvette. Mix gently and return

cuvette to ice. Spread 200 ml onto selective plates containing 1 M sorbitol.

EXAMPLE 4

Protocol: *E. Coli* Transformation

1. Thaw aliquot of competent subcloning efficiency HB101 (-70°C storage) on ice, keeping all reagents on ice. Use DH5a cells for lac z complementation; DM-1 for non-methylation. 2. Dispense 50 µl HB101 to required # of eppendorf tubes, one for each DNA sample, one for pBR322 (positive control) (pUC19 for DH5a and DM-1), one for no DNA (negative control). 3. Aliquot unused cells into 50 mL portions and refreeze in a dry ice/EtOH bath; store at-70°C. 4. Add 1 µl DNA (1 µg) to eppendorfs (1 µl of pBR322 or pUC19), tap tubes to mix, then incubate on ice for 30 min. For ligations, use 1-2 µl of ligation mixture (too much sours the transformation). 5. Heat shock at 37°C for 20 seconds. (45 sec at 42°C for DM-1 cells). 6. Place on ice for 2 minutes, then add 0.95 mL of room temp SOC media. Incubate at 37°C for 1 hour in bath (shaking optional) or on shaker in 37°C room. 7. Plate 100-200 µL of cells onto LB, 100 µg/mL Ampicillin, and incubate overnight at 37°C.

EXAMPLE 5

Protocol: GELase DNA Purification

Wizard PCR prep is an alternative protocol for DNA purification from a gel. GELase is recommended if the Wizard prep yield is low. Low yields happen if the desired fragments are less

than 200 kb or more than 5 kb long. 1. Separate DNA fragments on a 1% low melting agarose gel in fresh 1x TAE buffer. 2. Stain the gel with ethidium bromide in water. Using the hand-held UV lamp, cut out the fragments of interest with a new razor blade. 3. Place the gel slice in a pre-weighed eppendorf tube. Weigh both again to determine weight of gel slice. If the gel slice weight is more than 300 mg, split samples into two tubes after step 6. 4. Add 2 µl of 50x GELase buffer per 100 mg of gel slice. 5. Incubate the tube containing the gel slice at 70°C until the gel is completely molten. This will take at least 20 minutes. A good technique is to wait 30 minutes, pipette the mixture up and down a couple of times, then wait another 10 minutes. Be sure gel is completely melted. 6. Equilibrate the molten gel at 45°C for at least 30 minutes. 7. Add 1 U of GELase per 150 mg of molten agarose. Incubate for 4 hours. For >600 mg add 2 U of GELase. 8. Add 1 volume (1 volume = mg of gel slice) 5 M ammonium acetate to the solution. If using >300 mg of gel, a larger tube is needed for the ethanol precipitation. 9. Add 2 volumes (1 volume = mg of gel slice + ammonium acetate) of room temperature 100% ethanol and invert several times. 10. Pellet the DNA by centrifuging for at least 30 min at room temperature in 19 RAL. If the DNA concentration is very low, wait for 30 minutes after adding the ethanol and then centrifuge for 30 minutes. 11. Remove supernatant with a pipette and discard. 12. Wash the pellet with room temperature 70% ethanol. 13. Dissolve the DNA in water or TE. DNA can then be stored at -20°C.

EXAMPLE 6

Protocol: Ligation

Materials: T4 DNA ligase and 2x T4 DNA ligase buffer

For phosphatasing: Calf intestine phosphatase (CIP) and buffer (if needed). For blunt ending: dNTP mix (0.5 mM). Klenow fragment of E. Coli DNA polymerase I or T4 DNA polymerase. For linking: Oligonucleotide linkers- 0.2 mM DTT.

1. In a 20 µl reaction mixture, cleave the individual DNA components with appropriate restriction endonuclease. After the reaction is complete, inactivate the enzymes by heating 15 minutes to 65° C. If no further enzymatic treatments are necessary, proceed to step 6.

2. If the 5' phosphates of one of the DNAs are to be removed, add 2 µl of 10x CIP buffer and 1 U CIP; incubate 30 to 60 minutes at 37° C. After the reaction is complete, inactivate CIP by heating 15 minutes to 75° C. If no further enzymatic treatments are necessary, proceed to step 6.

3. For blunt ending, add 1 µl of a solution containing all 4 dNTPs (0.5 mM each) and an appropriate amount of the Klenow fragment of E. Coli DNA polymerase I or T4 DNA polymerase; carry out the filling in or trimming reaction. After complete, inactivate the enzymes by heating 15 minutes to 75°C. If oligonucleotide linkers are to be added, proceed to step 4. If a DNA fragment containing only one blunt end is desired, cleave the reaction products with an appropriate restriction endonuclease. If no further enzymatic treatments are necessary, proceed to step 6.

4. Add 0.1 to 1.0 µg of an appropriate oligonucleotide linker, 1 µl of 10 mM ATP, 1 µl of 0.2 M DTT, and 20 to 100 cohesive

- end units of T4 DNA ligase; incubate overnight at 15°C. Inactivate the ligase by heating 15 minutes to 75°C.

5. Cleave the products from step 4 with the restriction enzyme recognizing the oligonucleotide linker, adjusting the buffer conditions if necessary. If only one of the two ends is to contain a linker, cleave the products with an additional restriction enzyme.

6. Isolate the desired DNA segments by gel electrophoresis, if necessary. Then purify (GeneClean II or GelASE).

7. Ligation: 9 µl component DNAs (0.1 to 5 mg), 4 µl 5x ligase buffer, 1 mL (cohesive end) T4 DNA ligase (BRL: 1 unit = 300 cohesive end units, want 20 to 500 cohesive end units) water to 20 mL. Incubate 1 to 24 hours at 16°C.

8. Introduce 1 µl of the ligated products into competent E. coli cells and select for transformants. Then do miniprep and restriction mapping to screen for desired product.

EXAMPLE 7

Cloning:

All transformations were into *E. coli* strain DH5(following the manufacturer's protocol.

PCR

- Ampliwax PCR gem-mediated hot start PCR (Perkin-Elmer-Cetus, Norwalk, CT) - manufacturer's protocol for thin-walled tubes.
- GeneAmp PCR Core Reagents (Perkin-Elmer-Cetus)

• DNA Thermal Cycler 480 (Perkin-Elmer-Cetus)

AGA2

Cloned by PCR.

• Template for PCR was CEN BANK *S. cerevisiae* genomic library (American Type Culture Collection, Rockville, MD).

• Primers: 5'-ATTAGAATTCCCTACTTCATACATTTTCAA-3' and 5'-ATTACTCGAGCTATTACTGCAGagcgtagtctggaacgtcgtatgggtaAAAAACAT ACTGTGTGTTTATGGG-3'.

• Thermal profile:

Denaturation 1 minute at 94°C

Annealing 2 minutes at 41°C (first 5 cycles), 2 minutes at 45°C (25 additional cycles)

Extension 25 seconds at 72°C

Final polishing step 10 minutes at 72°C

The PCR product was cloned into plasmid pCR-Script using the pCR-Script SK(+) Cloning Kit (Stratagene, La Jolla, CA) following the manufacturer's protocol. The 342 bp AGA2 fragment was excised with EcoRI and XhoI, purified on a 1% agarose gel (protocol 6.1.7.2) and subcloned into pCR-Script containing the CUP1 promoter as a KpnI/EcoRI fragment.

HA peptide

The HA peptide was inserted by cassette mutagenesis. Complementary oligonucleotide strands encoding the Factor Xa recognition sequence and HA epitope were synthesized with cohesive overhangs allowing ligation to the 3' XhoI site of the AGA2 clone

while at the same time destroying this site; a downstream SacI site in pCR-Script annealed and ligated to the CUP1-AGA2 construct in pCR-Script. The insert included a new XhoI site at the 3' end of the HA sequence. CUP1-AGA2-HA was excised as a KpnI/XhoI fragment, purified on a 1% agarose gel, and subcloned into yeast shuttle vector pRS314 (1) already containing the alpha factor terminator sequence, to form surface display vector pCT101. Oligo sequences: 5'-TCGACGATTGAAGGTAGATACCCATACGACGTTCCAGACTACGCTCTGCAGT AATAGATTATCCTCGAGCT-3' and 5'-CGAGGATAATCTATTACTG CAGAGCGTAGTCTGGAACGTCGTATGGGTATCTACCTTCAATCG-3'.

The GAL promoter was excised from vector YCplac22-GAL. 12 bp palindromic linkers with appropriate cohesive overhangs were first cloned into this vector to alter restriction sites at both ends: EcoRI -> KpnI (E/KLINK) and BamHI -> EcoRI (B/ELINK). The resulting KpnI/EcoRI fragment was cloned into pCT101 to form vector pCT201. Oligonucleotide sequences: E/LLINK 5'-AATTGGTACC-3'; B/ELINK 5'-GATCGAATTC-3'.

The 4-4-20 scFv was amplified by PCR as above:

- Template: 4-4-20 in GeneX vector (obtained from D. Kranz, UIUC Dept. of Biochem.)
- Primers: 5'-ggttggccaagctagcGACGTCGTTATGACTCAA-3' and 5'-ggccggccaactcgagctattacaagtcttcttcagaaataagcttttgttcTGAGGAGACGGTG ACTGA-3'
- Thermal profile:

Denaturation 1 minute at 94°C

Annealing 2 minutes at 40°C (first 5 cycles), 2 minutes at 48°C (30 additional cycles)

Extension 50 sec at 72°C

Final polishing step 10 minutes at 72°C

The PCR product was cloned into pCR-Script and subcloned into pCT201 as a NheI/XhoI fragment using methods as above, creating vector pCT202. Vector pCT302 was created by inserting a synthetic oligonucleotide (UIUC Biotechnology Center) encoding a $(Gly_4\text{-}Ser)_3$ linker in frame between the AGA2 and 4-4-20 open reading frames of pCT202.

AGA1

Amplified by PCR as above:

- Template: CEN BANK
- Primers: 5'-ATTAGAATTCAGCTAAAAAAACCAAAAAAT-3' and 5'-ATTACTCGAGctaTTAACTGAAAATTACATTGC-3'
- Thermal profile:

Denaturation 1 minute at 94°C

Annealing 2 minutes at 41°C (first 5 cycles), 2 minutes at 45°C (25 additional cycles)

Extension 2 minutes, 20 sec at 72°C

Final polishing step 10 minutes at 72°C

The PCR product was gel purified using the GELase kit. The KpnI/SstI fragment of pCT201 was cloned into vector pRS316. This was then digested with EcoRI and XhoI, the excised AGA2 fragment removed by gel purifying the remaining vector fragment, and ligated to purified AGA1 PCR product digested with EcoRI and XhoI. The resulting clone was pCT211. The KpnI/SstI fragment of

pCT211 was subsequently cloned into vector YIplac211 to form vector pIU211.

EXAMPLE 8

Expression in yeast

Yeast strain *S. cerevisiae* BJ5465 (a ura3-52 trp1 leu2D1 his3D200 pep4:HIS2 prb1D1.6R can1 GAL). This strain is a pep4 and prb1 mutant making it deficient in proteases. Three nutritional markers have been deleted and may be used for plasmid selection: URA3, TRP1, and LEU2. HIS3 has been deleted, but the PEP4 deletion is covered with a HIS marker.

Transformation:

Vector pIU211 was cut with BsiWI which occurs uniquely within the AGA1 sequence. Approximately 100 ng of this linearized vector and 200 ng pCT202 were transformed simultaneously into yeast strain BJ5465 by electroporation (protocol 2.6.1). Transformants were selected on SD-CAA plates.

Experimental induction conditions:

A single colony of yeast transformed with pIU211 and pCT202 was inoculated into 3 ml SD-CAA and grown ~24 hours at 30°C. Cell density at this point was ~10^7-10^8 cells/ml (i.e., OD_{600} ~1-3). Sufficient cells were collected by centrifugation to inoculate 3 ml SG-CAA to a starting OD_{600} of ~0.5. This culture was grown ~ 20 hours at 30°C.

EXAMPLE 9

Fluorescent Labeling of Yeast Cells

The following method was used for the fluorescent labeling of yeast cells: 1. Collect 0.2 OD-ml (at 600 nm) of cells following growth for 20 hours in SG-CAA by centrifuging for ~10-30 sec at 14,000g. 2. Wash cell pellet by resuspending in TBS and spinning down. 3. Resuspend pellet in appropriate volume of TBS to make 100 µl final volume for incubation. Add the following amounts of labeling reagents, as appropriate: 1 µl of 25 mg/ml FITC-dextran (MW 2,000,000) (Sigma); 1 µl 9E10 Mab ascites fluid (Babco) or 10 µl 9E10 at 100 µg/ml (Santa Cruz Biotechnology); 100 µl 12CA5 Mab (Boehringer-Mannheim) at 10 µg/ml in TBS. Mix by vortexing or pipeting up and down. 4. Incubate 1 hour at room temperature, mixing cells approximately every 20 min by flicking tube, vortexing, or pipeting. 5. Spin down the cells and resuspend in appropriate volume of TBS to make 100 µl final volume. Add secondary reagents as appropriate in the following amounts: 4 µl a-mouse-PE (Sigma); 2 µl a-mouse-FITC (Sigma); 1 µl FITC-dextran. 6. Incubate 30 min at room temp. 7. Spin down and wash as in step 2. 8. Resuspend pellet in ~100 µl 10 mM Tris base, pH 8.3 (for anything labeled with FITC) or TBS for microscopy. For flow cytometry, resuspend in 500 µl 10 mM Tris (final cell density ~10^6/ml or more). Samples need to be in 0.5 ml microcentrifuge tubes for flow cytometry. For experiments using biotin-fluorescein, cells were grown, induced, harvested, and labeled as described with 10 µM biotin-fluorescein in place of FITC-dextran as the primary label, and a mixture of 3 µg of streptavidin-PE and 1 µg of RED613-conjugated goat anti-mouse $F(ab')_2$ (Life Technologies, Grand Island, NY) as the secondary labeling reagents.

EXAMPLE 10

Confocal fluorescence microscopy.

Yeast containing plasmid-directing surface expression of the HA peptide (pCT201) or the scFv fusion (pCT202) were grown for 20 hr in medium containing 2% galactose as the only carbon source and subsequently labeled with mAb 9E10, followed by a secondary anti-mouse IgG-R-phycoerythrin (PE) conjugate and FITC-dextran, as described. The labeled cells were mounted on polylysine-coated slides in 90% glycerol mounting medium containing 1 mg/ml *p*-phenylenediamine as an anti-bleaching reagent and analyzed with a laser scanning confocal microscope (UIUC Beckman Institute Microscopy Suite) at a rate of 8 seconds with a 63x power objective. Images from DIC, red PE fluorescence, and green FITC fluorescence were collected.

EXAMPLE 11

Flow cytometric analysis and sorting.

Labeled yeast cell suspensions were analyzed on a Coulter Epics XL flow cytometer at the Flow Cytometry Center of the UIUC Biotechnology Center. Event rate was maintained near 500 cells/sec. The population was gated by light scatter to avoid

examination of clumped cells, and data for 100,000 events were collected. For initial cell sorting experiments, yeast carrying the pCT202 vector were mixed with the untransformed parent strain BJ5465 and sorted based upon FITC signal on a Coulter 753 cell sorting bench modified with CICERO sorting electronics (UIUC Flow Cytometry Center). Presort and sorted samples were plated on non-selective medium, then replica plated onto medium selective for the pCT202 vector. Purity was determined as the fraction of non-selective colonies which are viable on selective plates.

EXAMPLE 12

Quantitation of surface antibody expression level.

Cells bearing vector pCT202 and Quantum Simply Cellular beads (Sigma, St. Louis, MO) were labeled with FITC conjugated mAb 12CA5 (Boehringer Mannheim, Indianapolis, IN) at 10 (g/ml in TBS as described and analyzed on a Coulter Epics XL flow cytometer. Comparison of the fluorescence intensity of the yeast sample with the standard beads allowed determination of antibody binding capacity of the displaying yeast cells by linear regression using QuickCal for Quantum Simply Cellular (Sigma).

EXAMPLE 13

Kinetic analysis of antigen dissociation from cells displaying scFv.

Yeast cells bearing plasmid pCT202 were grown and labeled with anti-c-*myc* mAb 9E10 and FITC-dextran or biotin-

fluorescein, as described. A fraction of the labeled population was analyzed flow cytometrically to determine the initial level of fluorescence. Non-fluorescent competitor (5-aminofluorescein) was added to a final concentration of approximately 10 μM (~1000-fold excess) and the FITC or PE fluorescence of the c-*myc* positive cell population was followed as a function of time at room temperature (21-23°C) on a Coulter Epics XL. Data were fitted as an exponential decay. The probability that a polyvalent antigen is bound to the cell as a function of time is given by $P = 1 - (1 - e^{-kt})^N$, where N is the valency, k is the kinetic rate constant for dissociation, and t is time. For long times t, this reduces to $P = Ne^{-kt}$. Thus, extrapolation of data for long t to time zero yields P = N, or a fluorescence intensity of $F_{ext} = N F_0$, where F_{ext} is the extrapolated fluorescence at the time of competitor addition and F_0 is the actual initial fluorescence. The valency of the interaction of surface displayed scFv 4-4-20 and polyvalent FITC-dextran was therefore determined as the y-intercept of the curves in Fig. 11.

Binding to soluble fluorescein (FDS) was assayed by observing fluorescence quenching by whole cells displaying scFv. Cells were suspended at 2×10^7 cells/ml in TBS + 0.1% BSA in a quartz cuvette thermostatted at 23°C and titrated with FDS over a range of 0-7.5 nM. Fluorescence at 520 nm was observed with an SLM Aminco SPF-500 spectrofluorometer using 488 nm excitation. Control cells displaying an irrelevant scFv were titrated to obtain a slope for a two-parameter fit of an equilibrium binding model to the data, yielding equilibrium constants and effective scFv concentrations. Following the equilibrium titration, 5-

aminofluorescein was added to 1 μM and the change in fluorescence of the sample followed with time to determine k_{off} for FDS.

EXAMPLE 14

Mutagenesis of scFv gene

Approximately 100 ng of pCT302 were transformed in duplicate into *E. coli* strain XL1-Red (Stratagene, La Jolla, CA) according to the manufacturer's protocol. Following 1 hr induction in SOC medium, the two transformant groups were pooled and 1/2000 of the pool plated on LB medium containing 100 μg/ml ampicillin to determine transformation efficiency. 5 ml of liquid LB medium containing 50 μg/ml ampicillin plus 100 μg/ml carbenicillin (LB-AMP50-CARB100) were inoculated with the remainder of the transformants and grown overnight at 37°C (OD_{600} ~1.0). A sufficient volume of this culture was collected to inoculate 50 ml LB-AMP50-CARB100 to OD_{600} = 0.01 in a baffled shake flask and grown to OD_{600} ~1.0 - 1.1 at 37°C. Cells were collected by centrifugation and used to inoculate 200 ml LB-AMP50-CARB100 to OD_{600} = 0.001, and the culture was grown at 37°C to OD_{600} ~1.0. Plasmid DNA was isolated by the QIAGEN Maxiprep kit (QIAGEN, Santa Clarita, CA). The recovered DNA was retransformed into XL1-Red and the growth cycle repeated three times, yielding a final product subjected to approximately 90 generations of growth in the mutator strain.

EXAMPLE 15

Library expression and kinetic screen

50 μg of mutagenized pCT302 DNA were transformed into yeast strain EBY100 by the method of Gietz and Schiestl in ten separate reactions. The products were pooled, and 1/2000 of the total plated on selective medium to determine the total number of transformants. The remainder were inoculated into 50 ml of selective glucose medium, grown overnight at 30°C, passaged to OD_{600} = 0.1, and expanded 10-fold. Selective galactose medium (5 ml) was inoculated to OD_{600} = 0.5 and grown overnight at 30°C to OD_{600} = 1.0-2.0. Samples of 10^7 cells (1 OD_{600}-ml) were labeled with FITC-dextran as described. Following labeling, cells were resuspended in 10 μM 5-aminofluorescein and 9E10 mAb at room temperature for 20 min, at which time samples were rinsed with ice cold buffer to stop competitive dissociation of FITC-dextran and labeled with anti-mouse-PE secondary antibody as described. Samples were sorted on a Coulter 753 bench with a sort window as shown in Fig. 4 and event rate of 4000/sec. 6 x 10^7 cells were examined during sorting round 1 and the window was set to collect 0.2% of the population. The collected cells were regrown in glucose medium and switched to galactose as described prior to repeating the competition and sorting. A total of four rounds of sorting and amplification were performed. 4 x 10^7 cells were examined in round 2, and 2 x 10^7 cells in each of rounds 3 and 4. Rounds 1 and 2 were performed in enrichment mode to provide a high recovery of all positive clones, and rounds 3 and 4 were performed in purify mode to reject coincident negative cells and achieve larger enrichment

factors. The products of round 4 were plated to isolate individual clones.

EXAMPLE 16

Establishment of Fusion Display System

A gene coding for a peptide epitope tag fusion with a yeast cell wall protein has been constructed and surface expression of the epitope verified. This cell wall protein, a-agglutinin, is involved in cell-cell adhesion during yeast mating and therefore is exposed on the external cell surface as a receptor for agglutinins on cells of the opposite mating type. Trial mixing and sorting experiments were performed to determine the one-pass and two-pass purification yields and purity for affinity screening by flow cytometry.

EXAMPLE 17

Yeast mating agglutinins

In the yeast life cycle, haploid cells occur as one of two possible mating types, a or α. When an a haploid cell and an α haploid cell come into physical contact, they adhere to one another through strong, specific interactions between cell surface adhesion proteins called "agglutinins." Once bound in this fashion, the cells fuse to form a diploid cell. As a platform for antibody display on the yeast cell wall, polypeptide fusions to a subunit of a-agglutinin were

constructed. Since the physiological role of agglutinins is to display protein binding sites on the exterior of the cell for specific, high-affinity interaction with other proteins, artifactual steric hindrance from yeast cell wall components was minimal.

As a eucaryote, yeast possesses a secretory apparatus which is highly homologous to that of antibody-secreting B lymphocytes. As a result, artifactual clone-dependent inefficient secretion should be minimized with this host. Numerous studies have revealed striking homology between the yeast and mammalian secretory pathways, such that particular molecules can be exchanged without significant loss of function, both *in vivo* and *in vitro*. Expression of mouse BiP functionally replaces yeast BiP, whose expression is essential for growth (Normington, 1989). Expression of mammalian PDI functionally replaces yeast PDI, another essential yeast ER lumenal protein (Gunther et al., 1993). Given the extensive similarities between yeast and mammalian secretion, clonal variability in antibody expression due to misfolding should be substantially reduced in yeast, compared to bacterial hosts. In fact, folding, assembly, and secretion of active antibodies in yeast has been demonstrated previously (Wood et al., '85; Horwitz et al., '88).

Yeast **a**-agglutinin is synthesized as two subunits: Aga1p, which possesses a phosphatidyl-inositol-glycan tail for anchorage to the cell wall and Aga2p, which binds to α-agglutinin with high affinity ($K_D \approx 1$ nM) (Lipke & Kurjan, 1992). Aga1p and Aga2p are linked by intersubunit disulfide bonds, and Aga2p is released to the growth medium after incubation with reducing agents such as DTT. Although phosphatidyl-inositol-glycan tails are generally localized to

a membrane, substantial evidence has accumulated that Aga1p is linked to the fibrous glucan in the cell wall by transglycosylation (de Nobel & Lipke, 1994).

EXAMPLE 18

Fusion construction

In order to establish the feasibility of using agglutinin fusions to display polypeptides on the yeast cell surface, an "epitope tag" peptide was first genetically fused to Aga2p. It is straightforward to extend this approach to antibody fusions. Passage of scFv molecules through the yeast secretory pathway is efficient, since folding, assembly, and secretion of active IgG's and Fab's has been demonstrated in yeast (Horwitz et al., 1988; Wood et al., 1985).

The AGA2 gene was cloned by PCR from a yeast genomic library and subcloned into an expression vector containing the strong copper-inducible CUP1 promoter which allows 25-fold variation of expression level. Coding sequence for the influenza HA epitope tag (Tyr-Pro-Tyr-Asp-Val-Pro-Asp-Tyr-Ala) was fused to the 3' end of the AGA2 open reading frame, preceded by a Factor Xa site-specific protease cleavage site (Ile-Glu-Gly-Arg'). The DNA sequence of this construct is shown in Figure 7. Convenient restriction sites have been included for in-frame fusion of single-chain antibody genes.

Although the Aga2p-HA fusion is simply a construction intermediate towards the final Aga2p-HA-antibody fusion, it does provide a means for confirming that a fusion peptide is anchored and accessible on the cell surface in this system. Anchorage of the HA

peptide to the external cell wall by the fusion has been verified by immunofluorescent staining of whole unfixed cells with the 12CA5 mAb (Boehringer Mannheim, Indianapolis, IN), detected by flow cytometry and fluorescence microscopy (Figure 9). Since whole 12CA5 antibody molecules bind to the HA epitope without any disruptive biochemical treatment of the cell wall, the Aga2p is accessible to the cell exterior for macromolecular recognition.

As described previously, the Aga2p binding subunit is attached to the cell wall through disulfide bonds to Aga1p, which is covalently anchored to other cell wall components. Treatment with DTT abolishes labeling with 12CA5, indicating that the Aga2p-HA fusion is attached to the cell surface by disulfide bonds. The AGA1 gene was cloned by PCR and subcloned downstream of the GAL1 promoter. Expression of AGA1 was induced by switching to galactose growth media.

The HA epitope tag was included in antibody fusions, to enable double fluorescence labeling for both surface antibody levels and binding of fluorescently-labeled antigens. This approach decouples cell-to-cell variations in antibody expression level from single-cell measurements of antigen affinity. For example, indirect immunofluorescence with phycoerythrin-labeled secondary IgG against the α-HA monoclonal antibody provides a measure of surface antibody numbers, while fluorescein-labeled antigen bound to the antibodies provides a measure of binding affinity. Because $\approx 10^4$ copies of a-agglutinin are displayed per cell, stochastic effects on binding measurements are minimal. Since commercial flow cytometers can detect under 10^3 fluorophore molecules, signal-to-

noise ratio should not be problematic. The ratio of green fluorescence (fluorescein, i.e., antigen binding) to red fluorescence (phycoerythrin, i.e., antibody number) is proportional to the fraction of antibodies bound by antigen.

In order to test the separation factors possible by this method, cells expressing the HA epitope tag were mixed at defined ratios with wild-type cells. The mixture of cells was labeled with fluorescein-labeled α-HA IgG and the most highly fluorescent subpopulation was sorted by flow cytometry. The sorted fraction was recultured and the fraction of cells bearing the Aga2p-HA fusion were determined by replica plating for a genetic marker associated with the expression vector. From this information, the single-pass purification possible by this method was estimated.

EXAMPLE 19

Fusion of a single-chain anti-fluorescein antibody to a-agglutinin

An anti-fluorescein single chain antibody based on the monoclonal antibody 4-4-20 has been constructed and characterized (Bird, '88; Davis et al., 1991). This single chain antibody is known to fold stably and retain affinity comparable to Fab fragments. The gene for monoclonal antibody 4-4-20 was fused to the existing AGA2-HA fusion gene for expression on the yeast cell surface.

Monoclonal antibodies against fluorescein are a useful model system for physicochemical studies of antibody-hapten interaction. The kinetics of antibody-antigen binding (Kranz et al.,

1982), thermodynamic analysis of complexation (Herron et al., 1986), the role of electrostatic interactions (Omelyanenko et al., 1993), and site-directed mutagenesis of an antibody binding site (Denzin et al., 1993) have been studied using this system.

Functionality of the displayed 4-4-20 fusion was determined by binding to multiply fluorescein-labeled dextran (Sigma). Since antibody binding quenches fluorescein emissions, the detected fluorescein represents moieties tethered to dextran which is bound to yeast via 4-4-20 bound fluorescein.

EXAMPLE 20

Development of display scaffold

Yeast possesses two related cell surface receptors known as a- and a-agglutinin that function to mediate cell-cell adhesion between a and a haploid cells as a prelude to fusion to form the diploid (Lu, 1995). α-agglutinin has been shown to be covalently linked to cell wall glucan by the C-terminus (Lu, 1995; Schreuder, 1993), and a-agglutinin is believed to be anchored by a similar linkage (Lu, 1995). Fusion to the C-terminal portion of α-agglutinin has been used previously to anchor enzymes and viral antigens on the yeast surface (Schreuder, 1993).

As a model system for development of the yeast surface display library screening method, we have displayed a functional anti-fluorescein scFv and c-*myc* epitope tag on the cell wall of yeast by fusion to a-agglutinin, which unlike α-agglutinin is a two-subunit glycoprotein (Fig. 2). The 725 residue Aga1p subunit anchors the

assembly to the cell wall (Roy, 1991) via β-glucan covalent linkage (Lu, 1995); the 69 amino acid binding subunit Aga2p is linked to Aga1p by two disulfide bonds (Cappellaro, 1994). The native a-agglutinin binding activity is localized to the c-terminus of Aga2p (Cappellaro, 1994); thus, this represents a molecular domain with accessibility to extracellular macromolecules and a useful site for tethering proteins for display. A vector for displaying proteins as C-terminal fusions to Aga2p was constructed (Fig. 3).

EXAMPLE 21

Verification of expression and surface localization of scFv

Expression of the Aga2p-scFv fusion is directed by the inducible GAL1 promoter (Johnston, 1984). Growth of yeast on glucose medium allows essentially complete repression of transcription from the GAL1 promoter, an important consideration for avoiding counterselection against sequences which negatively influence host growth. Switching cells to medium containing galactose induces production of the Aga1p and Aga2p fusion gene products, which associate within the secretory pathway and are exported to the cell surface. Surface localization of the Aga2p-scFv fusion has been verified by confocal fluorescence microscopy and flow cytometry. Cells labeled simultaneously with an anti-c-*myc* mAb and fluorescein-conjugated dextran (FITC-dextran) were examined by laser scanning confocal microscopy (Fig. 8). Control cells bearing a vector which directs display of an irrelevant peptide

(i.e., a hemagglutinin (HA) epitope tag only) are not labeled by mAb specific for the c-*myc* epitope or FITC-dextran (Fig. 8A).

In contrast, cells bearing the surface display vector pCT202 expressing the Aga2p-scFv-c-*myc* fusion are co-labeled by both the anti c-*myc* antibody and FITC-dextran (Fig. 8B), demonstrating that the antigen binding site is accessible to very large macromolecules. Both of these strains are positively stained by mAb 12CA5 directed against the HA epitope tag. Accessibility of the fusion for binding to both an intact IgG (150 kDa) and a 2 x 10^6 Da dextran polymer indicates an absence of significant steric hindrance from cell wall components, a significant advantage relative to *E. coli* surface displayed proteins which are buried within a lipopolysaccharide layer that forms a barrier to macromolecular diffusion.

Two-color flow cytometric analysis of these yeast strains likewise demonstrates accessibly displayed scFv on the cell surface. Negative control and scFv displaying strains were labeled with the anti-c-*myc* mAb 9E10 and FITC-dextran simultaneously. Bivariate histograms demonstrate a linear relationship between the intensity of phycoerythrin fluorescence (level of mAb 9E10 binding) and FITC fluorescence (antigen binding) for the cell population carrying the 4-4-20 display plasmid, while the control population exhibits background fluorescence (Fig. 9A and B). The distribution of fluorescence intensity within the positive fraction illustrates the importance of correcting the antigen binding signal for cell-to-cell variability in the number of displayed fusions, as determined by epitope tag labeling.

Quantitation of the display efficiency by comparison of an scFv-displaying cell population with calibration standards of known antibody binding capacities yields an average value of greater than $3x10^4$ fusions per cell. Treatment of cells displaying the Aga2p-scFv fusion with dithiothreitol prior to labeling eliminated staining of the cell surface by both FITC-dextran and mAb 9E10 (Fig. 9C), consistent with adherence of the fusion protein to the cell surface by a specific disulfide bonding interaction between the recombinant Aga2p subunit and Aga1p. This property illustrates another important feature of the yeast display system - proteins can be simply released from the cell surface by reduction for further characterization.

To examine further the specificity of the 4-4-20/fluorescein interaction, a competitive dissociation assay was performed using a non-fluorescent analog of fluorescein, 5-aminofluorescein. Analysis of these data yields a monovalent dissociation rate constant (k_{off}) at 21°C of 3.7 x 10^{-3} sec^{-1} for FITC-Dextran, and 3.9 x 10^{-3} sec^{-1} for fluorescein-biotin. Extrapolation of the exponential fit to t = 0 sec shows that the average valency of the interaction of a FITC-dextran molecule with scFv is less than 1.5. Similar results were obtained using fluoresceinated inulin, fluorescein-conjugated bovine serum albumin, and fluorescein-biotin as the competitor, indicating that the labeling of cells by FITC-dextran or fluorescein-biotin is due to a specific interaction between the displayed fusion and the fluorescein moiety. Furthermore, dissociation kinetics of fluorescein disodium salt (FDS) from surface displayed 4-4-20 scFv matched those from yeast-produced soluble 4-4-20 scFv as observed by spectrofluorometry.

EXAMPLE 22

Enrichment of displaying cells by flow cytometric cell sorting

To determine the effectiveness of flow cytometric sorting with yeast surface display, mixtures of yeast bearing the surface display vector with those lacking the associated selectable marker were sorted and purities independently determined by replica plating. Significant enrichment factors (up to 600-fold) are obtained (Table I). Thus, rare clones may be selected from yeast displayed libraries by initially enriching positive cells at relaxed stringency and high yield to provide a smaller population which can then be subjected to several passes of more stringent sorting to isolate rare clones.

TABLE I

Cells displaying 4-4-20 can be selectively enriched by flow cytometry

Initial Fraction w/ pCT202	Replica Colonies	Sorted Fraction w/ pCT202	Enrichment Factor
14%	48/58	83%	6x

0.5%	87/92	95%	200x
!!--0. 1%*	34/58	59%	~600x

*Initial fraction was estimated by flow cytometry.

EXAMPLE 23

Isolation of mutant scFv with lower k_{off} from a yeast displayed mutagenized library

Selection of scFv genes randomly mutagenized by propagation in a "mutator" strain of *E. coli* has been described (Low, 1996). A library of ~5 x 10^5 4-4-20 scFv mutants created by propagation of the yeast surface display vector in such a strain was expressed in yeast. The pool of cells displaying the scFv library were subjected to kinetic selection by competition of FITC-dextran labeled cells with 5-aminofluorescein. c-*myc* positive cells exhibiting the highest ratio of FITC to PE fluorescence were collected by flow cytometric sorting (Fig. 10A), amplified by regrowth under fusion repressing conditions (glucose carbon source), induced for surface fusion display, and resorted. Cells demonstrating a substantially increased persistence time of labeling by FITC-dextran were dramatically enriched following three rounds of sorting and amplification (Fig. 10).

FITC-dextran dissociation kinetics for two individual clones selected from the scFv library differed by 2.9-fold compared to wild-type 4-4-20 scFv (Fig. 11). Rate constants for the mutants were 1.9 x 10^{-3} sec^{-1} (mutant 4M1.1) and 2.0 x 10^{-3} sec^{-1} (4M1.2)

at 23°C, compared to 5.6 x 10^{-3} sec^{-1} for wild-type; similar experiments yielded k_{off} values for fluorescein-biotin of 2.4 x 10^{-3} sec^{-1}, 2.8 x 10^{-3} sec^{-1}, and 5.0 x 10^{-3} sec^{-1}, respectively. Additionally, soluble fluorescein dissociation kinetics determined by spectrofluorometry demonstrated a 2.2-fold improvement for both mutants relative to wild-type, and initial equilibrium fluorescence quenching experiments suggest a similar improvement in the affinity constant of the binding reaction. Isolation of clones with only threefold reduced off-rate demonstrates the capability of this screening method to achieve precise quantitative distinctions.

Of 26 selected clones individually analyzed, two were identically improved in k_{off} (4M1.1 and 4M1.2, described above); two demonstrated wild-type k_{off} with a decrease in c-*myc* labeling skewing the linear expression level/activity relationship; one exhibited wild-type k_{off} and c-*myc* labeling; and 21 bound with an apparent k_{off} approximately 10-fold lower than wild-type only to polyvalent 2 x 10^6 Da FITC-dextran, but not to monovalent FITC-dextran or fluorescein-biotin. Enrichment for clones with increased avidity resulted from use of polyvalent antigen (approximately 90 fluoresceins per dextran); avidity effects can be effectively avoided by appropriate design of screening conditions to ensure monovalent antigen binding. Furthermore, selection of epitope tag mutants can be eliminated by alternately detecting expression level by c-*myc* and HA tag labeling in sequential sorting rounds, or by alternative mutagenesis strategies targeting changes only to the scFv gene.

These results show that scFv fragments can be displayed on the surface of yeast in a manner accessible for macromolecular

recognition and amenable to combinatorial library construction and screening. The displayed scFv specifically binds antigen--the first demonstration of a functional antibody fragment displayed on the yeast cell surface. The application of this display system to library methods for *in vitro* antibody affinity maturation and for display of other mammalian proteins is a significant complementary alternative to existing technologies such as phage display, bacterial surface display, and the yeast two-hybrid method. Indeed, the literal first-attempt success of the yeast display system in recovery of improved fluorescein-binding scFv mutants from a relatively small library under non-optimized screening conditions clearly demonstrates the robustness of this technology. The demonstrated highly quantitative kinetic analysis of surface-tethered scFv and fine discrimination of clones with similar binding characteristics further attests to the great potential of yeast display for combinatorial optimization of proteins.

EXAMPLE 24

Display of an antibody to the T cell receptor in the yeast display system

Herein, a scFv (KJ16) specific for the Vb8 region of the T cell receptor (Roehm et al., 1985) was expressed in the yeast display system. This scFv-KJ16 inhibited the activity of T cells by competitively blocking the recognition of a TCR ligand such as the superantigen staphylococcal enterotoxin B (Cho et al., 1995). Since the affinity variants of this scFv may show enhanced T cell inhibition,

the use of the yeast display system in engineering higher affinity forms of scFv-KJ16 were examined.

A screen based on equilibrium binding of cell surface scFv to fluorescently-labeled antigen, a Vb8 single-chain TCR (Schlueter et al., 1996) was developed. Using two-channel flow sorting, selection was also based on the binding of a fluorescently-labeled anti-c-*myc* antibody to a ten-residue c-*myc* tag at the carboxy-terminus of the scFv. Variant scFv with a higher affinity for the TCR or with a lower affinity for the anti-c-*myc* antibody were isolated. As expected, the former had a mutation in a CDR (V_L CDR1) and the latter had a mutation in the c-*myc* epitope. Thus, these findings demonstrate that the yeast display approach can be used either to isolate higher affinity scFv or to identify the epitopes of a displayed protein recognized by a particular Mab.

Plasmids and Strains

The scFv-KJ16 V_L and V_H genes joined by a modified 205 linker (Cho et al., 1995) were subcloned by PCR into the vector pCR-Script (Stratagene, La Jolla, CA) following the manufacturer's protocol. A c-*myc* epitope tag was included at the carboxy-terminus of the scFv. The ~800-bp NheI/XhoI fragment containing the scFv was excised from pCR-Script and ligated into the yeast surface display vector pCT202 containing a nine-residue epitope tag (HA) and the AGA2 open reading frame downstream of the inducible GAL1 promoter. The resultant construct was transformed by the lithium acetate (LiAc) transformation method of Gietz and Schiestl (Gietz et al., 1995) into the *S. cerevisiae* strain BJ5465 (*a ura3-52 trp1 leu2D1*

his3D200 pep4::HIS2 prbD1.6 can1 GAL; Yeast Genetic Stock Center, Berkeley, CA) containing a chromosomally integrated AGA1 controlled by the GAL1 promoter (strain EBY100.

EXAMPLE 25

Induction and detection of scFv-KJ16 on the yeast surface

Yeast cells transformed with pCT202/scFv-KJ16 were grown overnight at 30°C with shaking in 3 ml selective glucose medium SD-CAA (glucose 2 wt %, Difco yeast nitrogen base 0.67 wt %, casamino acids 0.5 wt %). After ~18-20 hours, recombinant AGA1 + AGA2-scFv expression was induced at 20°C with shaking in 5 ml selective galactose medium (SG-CAA, where 2% galactose replaces the glucose in SD-CAA). Cultures were harvested after ~20-24 hours (1-2 doublings) by centrifugation, washed with PBS (10 mM $NaPO_4$, 150 mM NaCl, pH 7.3) containing 0.1% bovine serum albumin and 0.05% azide, and incubated 45 minutes on ice with 25 mL of 10 mg/ml anti-HA Mab 12CA5 (Boehringer Mannheim, Indianapolis, IN), anti-c-*myc* Mab 9E10 (1:100 dilution of raw ascites fluid; Berkeley Antibody Co., Richmond, CA), or biotinylated-scTCR [~360 nM] prepared from inclusion bodies expressed in *E. coli* (Schodin et al., 1996). Cells were washed with PBS and incubated 30 minutes on ice with either FITC-labeled $F(ab')_2$ goat anti-mouse IgG (1:50; Kirkegaard and Perry Labs, Inc., Gaithersburg, MD) or a streptavidin-phycoerythrin (SA-PE)

conjugate (1:100; PharMingen, San Diego, CA). Labeled yeast cells were analyzed on a Coulter Epics XL flow cytometer at the Flow Cytometry Center of the UIUC Biotechnology Center. Event rate was ~ 250 cells/sec. Data for 10,000 events was collected, and the population was gated according to light scatter (size) to prevent analysis of cell clumps. These conditions were also used to generate equilibrium antigen binding isotherms after incubation of scFv-KJ16 yeast with various dilutions of scTCR. Scatchard analysis was performed to determine the K_D values, using the estimated concentration of the biotinylated-scTCR and mean fluorescence units taken directly from flow data.

EXAMPLE 26

Production of a scFv-KJ16 random mutant library

Approximately 50 ng of pCT202/scFv-KJ16 were transformed into *E. coli* XL1-Red cells (Stratagene, La Jolla, CA) according to the manufacturer's protocol. Following a 1 hour induction in SOC medium, the recovery was centrifuged at 2000 rpm for 5 minutes and resuspended in 500 ml of liquid LB medium containing 100 mg/ml ampicillin plus 50 mg/ml carbenicillin (LB-AMP100-CARB50). The resuspension was added to 15-ml LB-AMP100-CARB50 in a 50-ml Erlenmeyer flask and grown at 37°C with shaking. The culture was replenished with a fresh 15-ml LB-AMP100-CARB50 at mid-log phase (OD_{600} ~ 0.2-0.4), then grown to saturation (OD_{600} ~ 1.0-1.1; this was considered one "cycle" or round of mutation). A small fraction of this culture (0.75 ml) was added to

the next cycle (15-ml LB-AMP100-CARB50). After six cycles of growth, Wizard Miniprep (Promega, Madison, WI) DNA plasmid preparations were performed on the 15-ml culture. Approximately 4.5 mg of pCT202/scFv-KJ16 DNA from cycle six were transformed into each of 3 tubes of yeast strain EBY100 using the LiAc method (Gietz et al., 1995). The 3 reactions were pooled and after resuspension in 1-ml ddH_2O, 1/2000 of the pool plated on selective plates to determine transformation efficiency. Fifty milliliters of SD-CAA were inoculated with the remainder of the culture, grown overnight at 30°C with shaking, passaged to OD_{600} = 0.05, and grown overnight at 30°C to OD_{600} >1.0. Five milliliters of SG-CAA were then inoculated to OD_{600} ~ 0.5 and grown overnight at 30°C with shaking to OD_{600} = 1.0-2.0.

EXAMPLE 27

Selection of scFv-KJ16 mutant library by FACS

Cells were double-labeled as described above with anti-c-*myc* Mab and biotinylated-scTCR (used at a concentration ~10 nM). The reaction volume was adjusted to maintain ~10-fold molar excess of antigen (scTCR) over surface scFv. Samples were sorted on a Coulter 753 bench with a sort window as shown in Figure 3 and an event rate of 4,000 cells/sec. A total of 8×10^7 cells were examined during the first sorting round, with 0.1-0.4% of the population collected. The collected cells were regrown at 30°C in SD-CAA and switched to SG-CAA prior to the next round of sorting. A total of 4 rounds of sorting was performed, with the first 2 sorts in enrichment

mode (high recovery of all positive clones) and the last 2 sorts in purification mode (coincident negative cells rejected). Immediately following the last sort, the collected cells were re-sorted and plated on selective plates to isolate individual clones.

EXAMPLE 28

Rescue and sequencing of mutant scFv-KJ16 genes

Plasmids from scFv-KJ16 yeast (wt and 2 mutants) were rescued according to the protocol described by Ward (Ward, 1991), except that cells were disrupted with a bead beater (BioSpec Products, Inc., Bartlesville, OK) for 2 minutes instead of vortexing. Cells were centrifuged for 1 minute and the upper (aqueous) phase collected. A Wizard Miniprep kit (Promega, Madison, WI) was used to prepare the plasmid DNA and *E. coli* DH5a competent cells (GibcoBRL, Gaithersburg, MD) were transformed with 1 ml of the DNA preparation using the $CaCl_2$ method. Transformations were plated on LB-AMP50. Sequencing of wt scFv-KJ16 and two mutants (mut4 and mut7) was performed using primers that flank the scFv of the display vector and fluorescence automated sequencing (Genetic Engineering Facility of the UIUC Biotechnology Center).

EXAMPLE 29

TCR Binding by Yeast Cell Surface scFv

The monoclonal anti-TCR antibody KJ16 recognizes a conformational epitope on the Vb8 chain of the TCR (Brodnicki et al., 1996). KJ16 has been used for many *in vivo* studies in mice, including efforts to target and delete the Vb8 population of T cells (Born et al., 1987, McDuffie et al., 1986, Roehm et al., 1985). To evaluate the possible effects of varying antibody affinity in mediating these effects, the use of a yeast display system to identify KJ16 variants with increased affinity for TCR was examined. The scFv gene from the anti-TCR antibody KJ16 has been cloned previously and the scFv protein exhibited approximately the same affinity, K_D ~ 120 nM, as KJ16 Fab fragments (Cho et al., 1995).

The scFv-KJ16 coding sequence was subcloned so as to be expressed as a fusion polypeptide with the Aga2p agglutinin subunit expressed on the yeast cell surface. The fusion polypeptide includes a hemagglutinin (HA) epitope tag N terminal to the scFv and a c-*myc* epitope tag at the carboxy-terminus. The inclusion of these epitopes allows monoclonal anti-HA (12CA5) and anti-c-*myc* (9E10) antibodies to be used in flow cytometry to quantify surface expression of the full length scFv independently of antigen-binding activity. Such normalization helps account for the effects of cell-to-cell variability in surface levels of the fusion polypeptide. As discussed below, the availability of two independent epitope tags can also control for the selection of individual epitope mutants that might not be desired in screening for ligand binding mutants. To evaluate the binding properties of cell surface scFv, a soluble single-chain Vb8-Va3 TCR (Schodin et al., 1996) was biotinylated and the bound ligand was detected with a phycoerythrin-streptavidin conjugate.

Figure 12 shows that yeast transformed with the scFv-KJ16/Aga2 plasmid expressed the HA epitope (Figure 12A) and the c-*myc* epitope (Figure 12B). Control yeast transfected with only the Aga2p/HA expression vector were positive for the anti-HA Mab but not for the anti-c-*myc* antibody. The fraction of cells in the non-fluorescent population has been found to depend on plasmid stability and culture growth phase (data not shown), but the physiological processes that are involved are unknown. Nevertheless, decreasing the induction temperature to 20°C and decreasing the induction time to less than two culture doublings produces populations with >75% of the cells displaying the scFv-KJ16. scFv-4-4-20 was displayed with this system with approximately the same proportion of positive cells.

Binding of biotinylated scTCR to cell surface scFv was also detected by flow cytometry (Figure 12C). The fraction of cells that expressed active scFv was similar to that detected with anti-HA and c-*myc* antibodies, consistent with the expression of full length, properly folded scFv. Furthermore, two-color histograms demonstrated a tight correlation of scTCR binding with both HA and c-*myc* epitope display (data not shown). Biotinylated-scTCR binding is specific to yeast displaying the scFv-KJ16, and was completely inhibited by excess soluble KJ16 IgG (Figure 12D).

The approximate affinity of the surface displayed scFv-KJ16 was determined *in situ* on the cell wall by titrating whole cells with varying concentrations of biotinylated scTCR. Equilibrium binding was measured by analyzing cell-bound scTCR by flow cytometry. Scatchard analysis of the binding data (Figure 13) yielded a K_D of 500 nM, within five fold of that observed for soluble scFv-

KJ16. Such agreement is reasonable, since K_D was calculated under the assumption that 100% of the scTCR was active, likely to be an overestimate (i.e. if only 20% were correctly folded, then the surface scFv would have a K_D ~ 100 nM). Previously, a substantial fraction of the scTCR purified from solubilized *E. coli* inclusion bodies is incorrectly folded was found (Schodin et al., 1996).

EXAMPLE 30

Selection of Mutagenized scFv-KJ16/Yeast by Fluorescence-Activated Cell Sorting

An *E. coli* mutator strain has been used to mutagenize an scFv for affinity maturation by phage display (Low et al., 1996). This approach was successful in identifying a mutant of scFv-4-4-20 with higher affinity for fluorescein using yeast display. A strength of this mutagenesis approach is its simplicity, requiring only *E. coli* transformation and cell growth. Furthermore, the *E. coli* mutator strain introduces mutations throughout the expression plasmid, and therefore does not bias changes to portions of the scFv believed to be important for determining binding characteristics. Whether this aspect of mutator strain mutagenesis is advantageous depends on the ability to identify key residues that might influence antigen binding, based on available structural information. Examination of published affinity maturation studies suggest that the location of such residues, generally in non-contact residues, is not yet predictable *a priori* (Hawkins et al., 1993, Patten et al., 1996, Schier et al., 1996, Thompson et al., 1996, Yang et al., 1995, Yelton et al., 1995).

To apply this strategy to scFv-KJ16, the scFv-KJ16/Aga2 plasmid was propagated in the *E. coli* mutator strain XL1-Red (Stratagene) for six cycles of growth. This procedure was predicted to introduce an average of two to three point mutations in the scFv coding sequence, based on a mutation rate per cycle of 1 in 2000 bps. The resultant plasmid preparation was transformed into yeast yielding a library size of approximately 3×10^5 transformants. In other work, larger libraries (10^7) have been obtained by further optimization of transformation procedures and by pooling independent transformations. This number does not represent an upper size limit for library construction, as further efforts at optimization and scaleup could be straightforwardly applied.

The mutagenized yeast library was subjected to four successive cycles of sorting and amplification, using a double stain for anti-c-*myc* antibody binding (FITC) and biotinylated-scTCR binding (PE). Biotinylated TCR was used at a 1:5000 dilution (~10 nM) that yielded just below the detectable threshold of binding by wt scFv-KJ16/yeast (Figure 13). The two channel fluorescence profiles of the mutated scFv-KJ16 sample after one sorting cycle (Figure 14A) and after four sorting cycles (Figure14B) are shown. Cells that exhibited fluorescence above the diagonal window shown in Figure 14 were collected for regrowth. The rationale for this diagonal window was that in any given round the sort criteria were based on antigen binding per displayed polypeptide fusion. For example, selection based only on higher PE fluorescence levels (i.e. scTCR binding) would include not only those mutants with higher affinity scFv, but those that display a higher density of scFv per yeast cell. The latter

mutants would in principle be eliminated by including the anti-c-*myc* antibody as one of the two parameters to normalize for surface expression variability. The first two sorting rounds were performed in enrichment mode, isolating the ~0.5% of the cell population with the highest fluorescence and not setting the sort software to reject coincidences (two cells in the same sorted droplet). The final two sorting rounds were performed for purity with high coincidence rejection. After the fourth cycle, cells were resorted immediately and plated. Ten colonies (mut1 - 10) were selected for further analysis.

EXAMPLE 31

Characterization of Mutant scFv-Yeast

Each of the 10 selected mutants were labeled with anti-HA antibody, anti-c-*myc* antibody, and biotinylated-TCR and was analyzed by flow cytometry (Figure 15). As might be expected, one clone (mut6) appeared phenotypically similar to wt scFv-KJ16/yeast. Another clone (mut7) was found to exhibit higher TCR binding levels, a result confirmed by several independent titrations. Finally, a number of the mutants (mut1-5, 8, 9) consistently showed reduced binding to the anti-c-*myc* antibody compared to binding of the anti-HA antibody or the biotinylated scTCR. The presence of this class of mutants could be explained by the diagonal sort window specification: as shown in Figure 14, cells can "move" into the sort window either by increasing scTCR (PE) binding at constant c-*myc* (FITC) signal, or alternatively by decreasing c-*myc* (FITC) binding at constant scTCR (PE) signal. The selection of these mutants could be easily

circumvented by using both epitope tags in the fusion, HA and c-*myc*. Thus, by alternating labeling of each of these epitope tags in each round of sorting, diminished binding to one of the epitope tags would not be enriched in consecutive sorting rounds as in this case.

Fluorescence histograms of the presumptive c-*myc* epitope mutant (mut4), the scTCR binding mutant (mut7) and another mutant (mut10) were compared with the wt scFv (Figure 16). Mut4 (Figures 16A and 16B) showed a reduction in anti-c-*myc* labeling, mut7 showed enhanced scTCR binding (Figures 16C and 16D), and mut10 did not show a shift in either, but the fraction of cells that were positive was higher than with the wt scFv (Figures 16E and 16F). As shown in Figures 16E and 16F, close to 100% of mut10 cells were positive for each of the agents tested. This contrasts with each of the other mutants (e.g. see mut4 and mut7) which resembled the wt scFv-KJ16 yeast in exhibiting two distinct populations of cells, one with reduced levels of cell surface scFv. Enhanced plasmid stability of mut10 and repeated failures to rescue the expression plasmid from mut10 into *E. coli* suggest that chromosomal integration has occurred with this mutant plasmid. Thus, the altered surface expression characteristics of mut10 appear to be a consequence of integration of the expression plasmid.

Binding affinity to scTCR was estimated for the mutants shown in Figure 16 by titration with soluble biotinylated scTCR (Figure 17). Nonlinear curve fitting of this data indicate unaltered K_D for mut4 and mut10, but a threefold increased affinity for mut7. The increase in mean fluorescence of mut10 is due to the absence of a

nonfluorescent tail in the distribution rather than increased scTCR binding, as is evident in Figures 16E and 16F.

EXAMPLE 32

Sequences of Mutant scFv

The nucleotide sequences of the wt-scFv-KJ16 cloned into the yeast display plasmid, and mut4 and mut7 following rescue of the plasmids from yeast was determined (Figure 18). The wt scFv-KJ16 contained two silent changes from the originally published scFv sequence (Cho et al., 1995). These may have been introduced by PCR prior to cloning of the scFv into the yeast display plasmid. The mut4 sequence contained one mutation and mut7 contained two mutations. The only mutation in mut4 was present in the c-*myc* epitope (Lys to Glu), consistent with its reduced binding by anti-c-*myc* antibody as described above. Mut7 contained a change from Arg to Lys in a framework region of the V_L region and a change from Ser to Arg in CDR1 of the V_L chain. The latter mutation is consistent with the higher binding affinity observed for mut7.

Discussion

Phage display has been used for the selection of scFv with higher antigen binding affinity, as well as isolation of new scFv's from naive libraries (Hoogenboom, 1997). However, there have been difficulties in the expression of some mammalian proteins in *E. coli*, in part because of toxicity, codon bias, or folding problems (e.g. Knappik & Pluckthun, 1995, Ulrich et al., 1995, Walker & Gilbert, 1994). Yeast expression can potentially obviate some of these problems, by

offering the advantage that proteins can be expressed with eucaryotic post-translational modifications (e.g., glycosylation and efficient disulfide isomerization). Furthermore, phage display does not generally possess the quantitative precision to discriminate between mutants with binding affinity differing by less than five-fold (Kretzschmar et al., 1995). By contrast, fluorescence labeling and sorting allowed the isolation of 4-4-20 scFv clones with only 3 fold increased affinity. Since most large changes in antigen binding affinity result from directed combination of point mutations, each with smaller effects (Hawkins et al., 1993, Schier et al., 1996, Yang et al., 1995), the capability to identify subtle improvements in affinity could be of significant value. With these advantages in mind, the use of a yeast display system for the affinity maturation of an anti-T cell receptor scFv was developed.

A scFv that is specific for the Vbs region of a mouse TCR was used in order to generate anti-TCR reagents that may ultimately have enhanced T cell targeting properties *in vivo* (Cho et al., 1997, Cho et al., 1995). The active scFv was expressed as an Aga2p fusion protein on the surface of yeast, with an affinity that was similar to the native scFv (~500 nM compared to 120 nM for the scFv). To select higher affinity scFv, random mutagenesis with a DNA-repair deficient strain of *E. coli* yielded a mutation frequency of ~2 to 3 per 1000 base pairs after six growth cycles. Flow cytometry with fluorescently labeled scTCR and anti-c-*myc* antibodies was used to sort cells displaying scFv's with increased scTCR affinity. The anti-c-*myc* antibody was included as a second criteria for selection to control for mutants with increased TCR binding due not to higher affinity but

because of higher cell surface expression of the scFv-c-*myc* fusion. After multiple rounds of selection, three mutant phenotypic classes were observed: 1) reduced binding to the c-*myc* antibody but unaltered scTCR binding (mut1-5, 8. 9); 2) enhanced binding to the scTCR with unaltered c-*myc* labeling (mut7); and 3) higher efficiency surface expression due to chromosomal vector integration (mut10).

The isolation of classes of mutants that are represented by mut4 and mut7 could be predicted from the selection criteria illustrated in Figure 14. That is, any mutant cell that was identified above the diagonal sort window boundary could be accounted for by either of the properties described for mut4 and mut7, since either an increase in scTCR (PE) signal or a decrease in c-*myc* (FITC) signal places a cell in the sorting window. This does not represent a substantial problem for this approach, however, because of the availability of two independent epitope tags. By utilizing the HA and c-*myc* tags in alternating sorting cycles, progressive enrichment for diminished labeling of one of the epitope tags should not occur.

The isolation of epitope tag mutants highlights an additional application for yeast surface display: mapping of epitopes recognized by monoclonal antibodies. Although alternative strategies that use peptide libraries have been successful in this regard for linear epitopes (Daniels & Lane, 1996), the approach described here can be extended to conformational epitopes. Accordingly, a properly folded protein can be displayed on the yeast cell surface and straightforward random mutagenesis as described herein can be applied to identify epitope residues from non-contiguous polypeptide sequence. Since nonfolded proteins are retained and degraded by the

eucaryotic secretory quality control apparatus and varied expression levels are identified by HA or c-*myc* labeling. false identification of epitope residues should be minimized by this procedure. The described approach is substantially easier than alanine scanning mutagenesis.

It is not clear why mut10 was enriched in this screen, since its average single chain T cell receptor labeling per c-*myc* labeling was unaltered. It is possible that the higher fraction of positively labeled cells biased this clone for enrichment due to random spillover into the sort window. In any case, neither scTCR or c-*myc* labeling were different for this clone, and structural rearrangements of the expression plasmid indicate that it had integrated into a chromosome.

The identification of a single unique CDR mutation in mut7 is consistent with the finding that this mutant scFv has enhanced binding to the T cell receptor. Future efforts to obtain only scFv with higher affinity for the T cell receptor (and not c-*myc* mutants) involves alternate selection with anti-HA and anti-c-*myc* antibodies to control for cell surface levels of the scFv. This strategy, combined with DNA shuffling techniques among selected mutants (Stemmer, 1994), should allow the isolation of scFv-KJ16 with considerably higher affinity than the wt scFv (K_D ~ 120 nM). Such mutant KJ16 scFv's can be used to test T cell signaling kinetic phenomena, as well as targeting of T cell-mediated killing via bi-specific antibodies (Cho et al., 1997, Rabinowitz et al., 1996).

The present invention demonstrates the purposeful isolation of affinity matured antibodies via cell surface display. As

described above, off-rate selection was employed to identify mutants with decreased dissociation rates, whereas in the expression of scFv-KJ16, equilibrated antigen binding was used. These two approaches are complementary, and depend on the affinity of the starting scFv. For K_D>1 nM, it is reasonable to pursue the strategy of equilibration with soluble labeled antigen as dissociation rates would be too rapid to allow effective discrimination of kinetic variation. Furthermore, at these lower affinities bulk soluble antigen is not substantially depleted from the labeling reaction mix. given that displayed scFv is present at effective concentrations of approximately 1-10 nM. By contrast, tightly binding antibodies such as 4-4-20 (K_D=0.4 nM) would deplete soluble labeled antigens at concentrations below K_D unless inconveniently large labeling volumes were employed. However, dissociation kinetics for such tightly binding antibodies are slow enough to enable quenching, sorting, and analysis via manual mixing procedures. Thus, one could employ a strategy whereby scFv's would be affinity matured via cycles of equilibrium-based screening and mutagenesis to reach K_D~1 nM, followed by cycles of off-rate screening and mutagenesis to obtain still further improvement.

Cell surface display and flow cytometric screening allows selection of clones from a library based on kinetic binding parameters such as K_D and the dissociation rate constant (k_{diss}). Binding parameters of selected mutants may then be quantitatively estimated *in situ* in the display format without a need for subcloning or soluble expression, as shown in Figure 17. By contrast, selection of phage displayed antibodies often involves increasingly stringent wash and elution conditions, even to the extent of pH 2 and 8 M GuHCl. Such

stringency selection has poor quantitative precision and may not always relate directly to binding parameters such as K_D or k_{diss} under ambient or physiological conditions.

Bacterial cell surface display systems have been described (Gunneriusson et al., 1996) for engineering of antibodies and other proteins. These systems possess some of the advantages of the present yeast display system, although they do not provide the post-translational processing capabilities of the eucaryotic secretory pathway. Access of macromolecules to the displayed protein on bacteria may also be restricted by the diffusion barrier presented by the lipopolysaccharide layer (Roberts, 1996). For this reason, binding to soluble protein antigens or epitope tag labeling with monoclonal antibodies is not possible. Surface display systems in cultured mammalian cells are also available (Rode et al., 1996), but construction and screening of combinatorial libraries for these systems are not as rapid or as versatile as for yeast.

A fairly small library (3×10^5) was screened to isolate the mutants described herein. This does not represent an upper limit on yeast library size. Yeast libraries with 10^7 clones have been constructed and further increases in library size, if necessary, would be attainable. The present invention shows that yeast surface display can be used to isolate a mutant scFv with increased affinity and that mutants with altered mAb epitopes can be enriched or excluded as desired. Further, the K_D can be estimated in situ in the display format without necessitating subcloning and soluble expression. Quantitative optimization of the screening conditions will enable further improvements in this method. Applications of yeast surface

display extend beyond antibody affinity maturation, to the isolation of binding domains from cDNA expression libraries, or isolation of mutant receptors or ligands on the direct basis of kinetic and equilibrium binding parameters.

EXAMPLE 33

Displayability and expression of the T cell receptor in the yeast display system

The present invention is also directed to a new process for engineering the T cell receptor for improved binding properties, e.g., to peptide-MHC complexes or superantigens. This invention establishes a method for displaying a T cell receptor in a yeast surface display library format. This method can be used to: 1) in general to express polypeptides that are not normally expressed on the surface or yeast, and 2) more specifically, to engineer higher affinity T cell receptors for a ligand of choice.

Protein engineering has not reached a level of development that allows rational and directed engineering of increased affinity binding. As a result, approaches have been developed that identify improved mutants from large mutant populations. The most widely used approach is "phage display", which has used to engineer antibodies, especially in the form of

linked, "single-chain" antibodies. However phage display methodology has been unable to display single-chain T cell receptors (scTCRs) successfully. This is most likely because folding of isolated single-chain T cell receptors is very inefficient in the absence of the other components of the CD3 complex and the protein folding machinery of the eucaryotic endoplasmic reticulum; the bacterial periplasm is unable to effectively fold these fragments.

The establishment of a yeast surface displayed T cell receptor is illustrated in Figures 19 through 21. A key improvement has been to isolate a mutant T cell receptor which can be displayed in this system. The wild-type T cell receptor is not functionally displayed, as shown by the absence of binding by an antibody (1B2) that is specific for the native conformation of the T cell receptor (Figure 19). By mutating the T cell receptor and screening a library for 1B2 binding, a mutant single chain T cell receptor displayed in yeast was identified. This establishes a system which can now be used to isolate mutant single chain T cell receptors with improved binding properties.

The present invention provides a yeast cell-surface display system successful in expressing the T cell receptor. Second, expression of the full length T cell receptor could only be achieved after randomly mutagenizing the T cell receptor gene and then selecting by flow cytometry for surface expression. This method thereby exploited an evolutionary approach to "correcting" the expression defect in the T cell receptor.

This same approach could be applied to any polypeptide which in its wild-type form is not displayed efficiently. Selection for

"displayability" has been reduced to practice for the T cell receptor, as described in examples 33-37. Once displayable mutant versions of the polypeptide are obtained, these versions can then be subjected to the screening processes for improved binding properties that are described in examples 1-32.

Improved T cell receptor molecules are useful in therapies for cancer, sepsis, and autoimmune diseases such as arthritis, diabetes, or multiple sclerosis. For example, soluble forms of high affinity T cell receptors would act as antagonists of detrimental T-cell mediated autoimmune diseases and thereby provide potential treatments for these diseases. Analogous strategies have been successfully employed with a soluble tumor necrosis factor receptor (TNF-R) and forms of this receptor are in clinical trials for septic shock and rheumatoid arthritis (Moosmayer et al., 1995).

In the methods of the present invention, yeast surface display allows single chain T cell receptors to be engineered to bind with high affinity to MHC-peptide complexes or superantigens. Such molecules would find a variety of medical uses. Examples include, but are not limited to: 1) interfering with inappropriate T cell attacks on healthy tissue in autoimmune diseases such as arthritis, diabetes, and multiple sclerosis; 2) interfering with septic shock due to bacterial superantigen that interact with T cells, leading to massive inflammatory reactions; and 3) destruction of tumor cells that bear T cell receptor ligands (e.g. specific tumor peptide/MHC complexes) by using high affinity T cell receptor together with anti-CD3 bispecific agents to redirect T cells to attack the cancerous cells.

Plasmids and strains. The single-chain TCR gene (V(8.2-linker-V(3.1) gene joined by a modified 205 linker (Cho et al., 1995) was subcloned by PCR into the vector pCR-Script (Stratagene, La Jolla, CA) following the manufacturer's protocol. A 6-His epitope tag was included at the carboxy-terminus of the scTCR for purification purposes. The ~800-bp NheI/XhoI fragment containing the scTCR was excised from pCR-Script and ligated into the yeast surface display vector pCT202 containing a nine-residue epitope tag (HA) and the AGA2 open reading frame downstream of the inducible GAL1 promoter. The resultant construct was transformed by the lithium acetate (LiAc) transformation method of Gietz and Schiestl (Gietz et al., 1995) into the *S. cerevisiae* strain BJ5465 (αura3-52 trp1 leu2Δ1 his3Δ200 pep4::HIS2 prbΔ1.6 can1 GAL; Yeast Genetic Stock Center, Berkeley, CA) containing a chromosomally integrated AGA1 controlled by the GAL1 promoter (strain EBY100).

EXAMPLE 34

Production of an scTCR random mutant library

Approximately 50 ng of pCT202/scTCR were transformed into E. coli XL1-Red cells (Stratagene, La Jolla, CA) according to the manufacturer's protocol. Following a 1 hour induction in SOC medium, the recovery was centrifuged at 2000 rpm for 5 min. and resuspended in 500 µl of liquid LB medium containing 100 µg/ml ampicillin plus 50 µg/ml carbenicillin (LB-AMP100-CARB50). The resuspension was added to 15-ml LB-AMP100-CARB50 in a 50-ml

Erlenmeyer flask and grown at 37°C with shaking. The culture was replenished with a fresh 15-ml LB-AMP100-CARB50 at mid-log phase (OD_{600} (0.2-0.4), then grown to saturation (OD_{600} ~1.0-1.1; this was considered one "cycle" or round of mutation). A small fraction of this culture (0.75 µl) was added to the next cycle (15-ml LB-AMP100-CARB50). After six cycles of growth, Wizard Miniprep (Promega, Madison, WI) DNA plasmid preparations were performed on the 15-ml culture. Approximately 10 µg of pCT202/scTCR DNA from cycle six were transformed into each of 10 tubes of yeast strain EBY100 using the LiAc method. The 10 reactions were pooled after resuspension in 1-ml ddH_2O/tube, 1/10,000 of the pool plated on selective plates to determine transformation efficiency. The library size was approximately 7 x 10^6. A 50 ml volume of SD-CAA (glucose 2 wt %, Difco yeast nitrogen base 0.67 wt %, casamino acids 0.5 wt %) was inoculated with the remainder of the culture, grown overnight at 30°C with shaking, passaged to OD_{600} = 0.05, and grown overnight at 30°C to OD_{600} >1.0. Five milliliters of selective galactose medium SG-CAA (where 2% galactose replaces the glucose in SD-CAA) were then inoculated to OD_{600} = 0.5 and grown overnight at 20°C with shaking for ~20-24 h (1-2 doublings).

EXAMPLE 35

Selection of scTCR mutant library by fluorescence-activated cell sorting

Cells were labeled with 25 µL Mab 1B2 (anti-Vβ8.2Vα3.1; prepared from ascites fluid and conjugated to biotin) at a

concentration of 20 µg/ml. Samples were sorted on a Coulter 753 bench with an event rate of ~4,000 cells/sec (Flow Cytometry Center, UIUC Biotechnology Center). A total of 6 x 10^7 cells were examined during the first sorting round, with ~5% of the population collected. The collected cells were regrown between sorts at 30°C in 4 ml selective glucose medium SD-CAA. After ~18-20 hours, recombinant AGA1 + AGA2-scFv expression was induced at 20°C with shaking in 5 ml SG-CAA. A total of 3 rounds of sorting was performed, with the first sort in enrichment mode (high recovery of all positive clones) and the last 2 sorts in purification mode (coincident negative cells rejected). Immediately following the last sort, the collected cells were re-sorted, collected as two separate populations ("high expression" and "low expression"), and plated on selective plates to isolate individual clones. Twenty clones were examined by flow cytometry.

EXAMPLE 36

Induction and detection of mutant scTCR on the yeast surface

Individual clones from the pCT2(2/scTCR library sorting were grown overnight at 30°C with shaking in 3 ml SD-CAA followed by induction in SG-CAA as described above. Cultures were harvested after (20-24 hours (1-2 doublings) by centrifugation, washed with PBS (10 mM $NaPO_4$, 150 mM NaCl, pH 7.3) containing 0.1% bovine serum albumin and 0.05% azide, and incubated 45 minutes on ice with 25 (L of 10 µg/ml anti-HA Mab 12CA5 (Boehringer Mannheim,

Indianapolis, IN), or biotinylated-1B2 Mab (20 µg/ml) prepared from ascites fluid. Cells were washed with PBS and incubated 30 minutes on ice with either FITC-labeled $F(ab')_2$ goat anti-mouse IgG (1:50; Kirkegaard and Perry Labs, Inc., Gaithersburg, MD) or a streptavidin-phycoerythrin (SA-PE) conjugate (1:100; PharMingen, San Diego, CA). Labeled yeast cells were analyzed on a Coulter Epics XL flow cytometer. Event rate was ~ 250 cells/sec. Data for 10,000 events was collected, and the population was gated according to light scatter (size) to prevent analysis of cell clumps. Results from the wild type (wt) TCR and several representative TCR mutants are shown in Figure 19. Double mutants containing the combined mutations from several of these isolates were also constructed and the results of flow cytometry of these are shown in Figure 20.

EXAMPLE 37

Rescue and sequencing of mutant scTCR genes

Plasmids from scTCR yeast (wt and 20 mutants) were rescued according to the protocol described by Ward (Ward, 1991), except that cells were disrupted with a bead beater (BioSpec Products, Inc., Bartlesville, OK) for 2 minutes instead of vortexing. Cells were centrifuged for 1 minute and the upper (aqueous) phase collected. A Wizard DNA Clean-Up kit (Promega, Madison, WI) was used to prepare the plasmid DNA and *E. coli* DH5α ElectroMAX competent cells (GibcoBRL, Gaithersburg, MD) were transformed via electroporation with 1 µl of the DNA preparation. Transformations were plated on LB-AMP50. Sequencing of wt scTCR and 20 mutants

(mTCR1-mTCR20) was performed using primers that flank the scTCR of the display vector and fluorescence automated sequencing (Genetic Engineering Facility of the UIUC Biotechnology Center). Single mutations were found in the TCR for each of the isolates shown (Figure 21). These mutations may allow the expression of more stable TCR for possible therapeutic uses.

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Any patents or publications mentioned in this specification are indicative of the levels of those skilled in the art to which the invention pertains. These references are incorporated by reference to the same extent as if each individual publication was specifically and individually incorporated by reference.

One skilled in the art will readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The present examples along with the methods, procedures,

molecules, and specific compounds described herein are presently representative of preferred embodiments, are exemplary, and are not limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention as defined by the scope of the claims.